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(54) Tear resistant multilayer films based on sebacic acid copolyesters and articles incorporating such films

Reissfeste Verbundfilme auf Basis von Copolyester der Sebacinsäure und solche Filme enthaltende Gegenstände

Films multicouches résistants à la déchirure à base de copolyesters d'acide sébacique et articles incorporant de tels films

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EP 0 591 055 B1

DescriptionField of the Invention

5 This invention relates to multilayer films, and, more particularly, to tear resistant multilayer films based on sebacic acid copolyesters.

Description of the Related Art

10 Traditionally, "tear resistance" has described the ability of a film to resist continuing to tear once a tear has been started. Trash and grocery bags, often based on polyolefins such as polyethylene, are examples of films that are conventionally considered to be tear resistant. These films have considerable stretch which enables them to resist advancing an already formed tear. By "stretch" it is meant that the films have a low tensile modulus and are not dimensionally stable.

15 Also known are films which are relatively stiff. In this regard, "stiff" refers to films which cannot be stretched significantly without breaking; that is, films which are dimensionally stable, creep-resistant (stretch resistant), and of high modulus. Examples of stiff, dimensionally stable, high modulus materials are certain packaging films such as cellophane, polyesters and biaxially oriented polypropylene. However, these films have low tear resistance. That is, once a tear has been started, the film continues to tear quite easily. resistance. That is, once a tear has been started, the film continues to tear quite easily.

20 There are numerous applications where stiff, tear resistant films would be desirable. For example, films which provide sign faces and building awnings must be tear resistant to have a useful life. On the other hand, these films must also be relatively stiff so that they will not billow in the wind or sag with age.

25 Backings for abrasive sanding belts experience harsh operating conditions and must resist tearing. However, sanding belts which stretch are undesirable because they may not fit securely on the sander and may work free under normal use.

Angioplasty balloons for expanding blood vessels during surgery require stiff, tear resistant films. The balloons cannot readily shatter (i.e., tear) during use. The balloons must also inflate to a controlled size and should not stretch to a larger size.

30 For certain tapes, stiff, tear resistant backings would be desirable. Such backings would not readily continue to tear if inadvertently nicked or cut when dispensed. At the same time, the backings would be stretch resistant which could enhance the stability of articles taped therewith.

Films for shatterproofing windows need to be tear resistant. However, the performance of such films would be enhanced if the films were also stiff and tear resistant as the combination of these properties would help the film to absorb energy in the event of a window shattering impact.

35 Numerous packaging films are disclosed in the prior art. U.S. Patent No. 3,188,265, "Packaging Films" issued June 8, 1965 to R. Charbonneau, et al. discloses a heat-sealable film comprising polyethylene extruded onto a web of oriented polyethylene terephthalate. U.S. Patent No. 4,705,707 "Polyethylene/Polyester Non Oriented Heat Sealable, Moisture Barrier Film and Bag," issued November 10, 1987 to J. Winter discloses a moisture barrier film useful in micro-waveable food pouches. The film comprises three and five layer nonoriented structures of polyethylenes and polyesters or copolyesters.

40 U.S. Patent No. 4,965,108 "Low Temperature Impact and Puncture Resistant Thermoplastic Films and Bags Therefrom," issued October 23, 1990 to E. Biel et al. discloses multilayer film and bag structures comprising a polypropylene copolymer inner layer, an outer layer (e.g., a polyester or a polyamide), and a polypropylene based bonding resin therebetween.

45 U.S. Patent No. 4,636,442 "Laminated Structures of Polyethylene Terephthalate and Elastomeric Copolyesterethers," issued January 13, 1987, to R. Beavers et al. discloses multilayer films reportedly having improved flex-crack resistance. The films are based on polyethylene terephthalate and elastomeric copolyesterethers. Biaxially oriented three and five layer films in which the amount of copolyesterether is from about 5 to about 75 weight % (preferably 10 to 60 weight %) are disclosed.

50 U.S. Patent No. 4,939,009 "Multilayer Sheets Having Excellent Adhesion," issued July 3, 1990, also to R. Beavers et al., discloses three and five layer films based on polyolefins and copolyesterethers with tie layers therebetween.

U.S. Patent No. 4,729,927 "Polyester Packaging Material," issued March 8, 1988, to M. Hirose et al. discloses a packaging material comprising polyethylene terephthalate and a second material based on polyethylene isophthalate copolymerized with an aliphatic hydroxycarboxylic acid having up to eight carbon atoms. Reportedly, the number of layers is not particularly critical, although films with up to five layers are said to be preferred.

Japanese Kokai Patent No. 2-270553 "Multilayer Plastic Sheet with Gas Barrier Feature," published November 5, 1990 discloses multilayer films based on layers of saponified ethylene/vinyl acetate copolymers, modified polyolefin

adhesives, and thermoplastic polyesters.

Impact resistant and/or shatterproof security films for windows are also known. For example, U.S. Patent No. 3,899,621 "Security Film for Shatter-Proofing Windows," issued August 12, 1975 to M. Willdorf discloses three and five layer films comprising layers of polyesters and polyurethanes. Preferably, the polyester layers range in thickness from 0.5 to 5 mils and the polyurethane layers range in thickness from 0.2 to 0.4 mil. U.S. Patent No. 3,891,486 "Process for Producing Solar Control Window," issued June 24, 1975, also to M. Willdorf, discloses a solar control film comprising a pair of polyester (e.g., polyethylene terephthalate) layers each from 0.25 to 1 mil thick with a vapor-deposited aluminum coating and an adhesive therebetween.

U.S. Patent No. 4,945,002 "Impact-Resisting Anti-Lacerative Window Units," issued July 31, 1990 to I. Tanuma et al. discloses a three layer film comprising two exterior layers (e.g., an ethylene/vinyl acetate copolymer, an ethylene/vinyl acetate/triallyl isocyanurate terpolymer, a polybutyl butyryl, a polyvinylformal, or a polyurethane), and an intermediate layer (e.g. polyethylene terephthalate, polyamides, polyester polyethers, polysulfones or polyimides) therebetween. The film is sandwiched between a pair of transparent glass or plastic plates.

Various tapes are also known. For example, U.S. Patent No. 4,091,150 "Coextruded Polyester Splicing Tape," issued May 23, 1978 to G. Roelofs discloses a multilayer tape comprising a support film formed from a tough, flexible polyester (e.g. polyethylene naphthalate or polyethylene terephthalate) which is coextruded with an adhesion promoting polyester. A thermoset adhesive is adherently bonded to the adhesion promoting polyester.

U.S. Patent No. 4,908,278 "Severable Multilayer Thermoplastic Film," issued March 13, 1990 to Bland et al. discloses a multilayer film which may be easily and precisely cut in a straight line. The film comprises alternating layers of brittle and ductile materials. Japanese Kokoku Patent Publication No. 63-5394 "Laminate Film," published October 26, 1988, discloses three and five layer tape backing films comprising layers of different polyesters. Reportedly, the films have good manual tearing properties.

U.S. Patent No. 4,540,623 "Coextruded Multi-layer Articles," issued September 10, 1985 to J. Im et al. discloses an impact resistant multilayer laminate comprising alternating layers (preferably at least about 40 layers) of coextruded polymeric thermoplastics wherein one of the materials contains a carbonate polymer. Suggested uses include glazing applications for windows and signs.

European Patent Application No. 0,426,636 "Iridescent Film with Thermoplastic Elastomeric Components," published May 8, 1991 discloses a transparent thermoplastic film of at least ten layers. The adjacent layers differ in refractive index and at least one of the layers is based on a thermoplastic elastomer resin. The layers range in thickness from 30 to 500 nanometers.

The article "Some Physical Properties of Multilayered Films" by W.J. Schrenk and T. Alfrey Jr., Polymer Engineering and Science, November, Vol.9, No.6, p. 393 - 399, 1969, discloses multilayered films comprising layers of high and low modulus materials.

The article "Advanced Polymers" by Eric Bauer, Scientific American, October, 1986, pages 161, 164, discloses a sheet of alternating layers of stiff and ductile materials.

SUMMARY OF THE INVENTION

In general, this invention relates to a tear resistant film comprising at least three layers situated one on the other in a parallel array. The layers occur essentially randomly in the array and are individually selected from a stiff polyester or copolyester and a ductile sebacic acid based copolyester. The film comprises at least a layer of a stiff polyester or copolyester and at least a layer of a ductile sebacic acid based copolyester. Preferably, the stiff polyester or copolyester layers are oriented in at least one direction and, more preferably, are biaxially oriented.

By "tear resistant" it is broadly meant that a film according to the invention demonstrates a Graves area in one direction of the film which exceeds the Graves area in the same direction for a single layer film comprising only the stiff polyester/copolyester of the multilayer film, the single layer film being processed in the same manner as and to substantially the same thickness as the multilayer film. Preferably, multilayer films according to the invention demonstrate a Graves area in one direction of the film equal to at least about $40 + 0.4(x)$ kpsi% wherein x is the nominal thickness of the film in microns. More specifically, Graves area is obtained by mathematically integrating the area beneath the curve in a graphical plot of the stress (as measured in kpsi) experienced by the film versus the strain (as measured by Graves elongation in % which is defined more fully below) that the film undergoes during a test in which a film sample specifically shaped for the Graves area test is clamped between opposed jaws that are moved apart at a constant rate to concentrate the tearing stresses in a small area. Thus, Graves area is a combined measure of the film's tensile modulus (i.e., the film's stiffness and dimensional stability) and the ability of the film to resist advancing a tear. Consequently, Graves area may be regarded as a measure of the total energy required to cause the film to fail; that is, the ability of the film to absorb energy.

Moreover, preferred multilayer films desirably exhibit a Graves elongation at break (defined below) of at least 20%, more preferably at least 40% during the Graves area test. In addition, preferred multilayer tear resistant films according

to the invention demonstrate a tensile modulus (as measured in a conventional tensile test) of at least 175 kpsi (1,208 MPa), more preferably at least 240 kpsi (1,656 MPa), and most preferably at least 450 kpsi (3,105 MPa) in at least one direction of the film.

Both the thickness of the film and the individual layers which comprise the film may vary over wide limits. Films according to the invention typically have a nominal thickness of from about 7 to 500 μm , more preferably from about 15 to 185 μm . The individual layers of stiff polyester or copolyester typically have an average nominal thickness of at least about 0.5 μm , more preferably from greater than 0.5 μm to 75 μm and, most preferably, from about 1 to 25 μm . It is preferred that the ductile sebacic acid based copolyester layers be thinner than the stiff polyester/copolyester layers. The ductile material layers may range in average nominal thickness from greater than about 0.01 μm to less than about 5 μm , more preferably from about 0.2 to 3 μm .

Similarly, the exact order of the individual layers is not critical. The total number of layers may also vary substantially. Preferably, the film comprises at least 3 layers, more preferably from 5 to 35 layers, and most preferably 13 layers.

Stiff polyesters and copolyesters according to the invention are typically high tensile modulus materials, preferably materials having a tensile modulus, at the temperature of interest, greater than 200 kpsi (1,380 MPa), and most preferably greater than 400 kpsi (2,760 MPa). Particularly preferred stiff polyesters and copolyesters for use in films according to the invention comprise the reaction product of a dicarboxylic acid component selected from the group consisting of terephthalic acid, naphthalene dicarboxylic acid and ester derivatives thereof, and a diol component selected from the group consisting of ethylene glycol and 1,4-butanediol. Additional stiff copolyesters based on these materials may also be provided by copolymerizing these ingredients with one or more other diacids and/or one or more other diols.

Ductile sebacic acid based copolyesters useful in the practice of the invention generally have a tensile modulus of less than 200 kpsi (1,380 MPa) and a tensile elongation (as defined below), at the temperature of interest, of greater than 50%, preferably greater than 150%. A preferred ductile copolyester comprises the reaction product of 20 to 80 (more preferably 70 to 50, and most preferably 60) mole equivalents terephthalic acid (or an ester derivative thereof), correspondingly, 80 to 20 (more preferably 30 to 50, and most preferably 40) mole equivalents sebacic acid (or an ester derivative thereof), and 100 mole equivalents ethylene glycol. The terephthalic acid may be replaced in whole or in part by naphthalene dicarboxylic acid such as dimethyl 2,6-naphthalene dicarboxylic acid (or an ester derivative thereof). In another preferred embodiment, a portion of the sebacic acid is replaced by an equivalent amount of cyclohexane dicarboxylic acid (or an ester derivative thereof).

Surprisingly, beneficial improvements in the tear resistance of films comprising alternating layers of stiff polyesters/copolyesters and ductile sebacic acid based copolyesters are realized when the ductile material provides less than 5 weight % of the film. Ductile material amounts of at least about 1 weight % (preferably at least about 2.6 weight %) up to about 10 to 20 weight % of the film may be used.

Films according to the invention may optionally include a layer of an intermediate material disposed between otherwise adjacent layers of the stiff and ductile polymers. Useful intermediate materials may be selected from a wide variety of polymers and, in some cases, may be selected to enhance the adhesion between the otherwise adjacent stiff and ductile layers. One or more functional layers may also be applied to one or both of the major surfaces of the film.

Multilayer films according to the invention provide an improved combination of stiffness and tear resistance especially when compared to films comprising only a single layer of one of the materials or single layer blends of both materials. Films according to the invention are useful in a wide variety of products, including, for example, sign faces and backings for coated abrasive articles.

The multilayer tear resistant films of the invention are particularly useful as security control laminates for shatterproofing glazing members against impact or explosion. In one embodiment of this application, the invention pertains to a security control laminate comprising a first tear resistant film having a first face and a first layer of adhesive on the first face for bonding the laminate to a glazing member. Typically, the adhesive coated face of the tear resistant film is temporarily disposed on a removable release liner which is discarded during application of the laminate to the glazing member. The security control laminate may further comprise means for absorbing ultraviolet radiation such as a coating layer interposed between the first tear resistant film and the layer of adhesive. Security control laminates according to the invention may also comprise a dyed film (bonded to the second face of the tear resistant film) and an abrasion resistant coating on the otherwise exposed surface of the dyed film.

In other embodiments, the security control laminate may comprise a second tear resistant film which is adhesively bonded to the first film. Such constructions may also include ultraviolet radiation absorbent and abrasion resistant coatings. Also contemplated is the inclusion of a metalized layer for imparting solar control properties to the security control laminate. A metalized layer may comprise an optically clear film having a layer of aluminum, gold, silver, copper, nickel and the like thereon. The security control laminate may be applied to a single glazing member or positioned between two glazing members. The glazing member(s) can be mounted within a frame to which the security control laminate may be optionally secured.

BRIEF DESCRIPTION OF THE DRAWINGS

The invention will be more fully understood with reference to the following drawings in which similar reference numerals designate like or analogous components throughout and in which:

- FIG. 1 is an enlarged perspective view of a multilayer tear resistant film according to the invention;
 FIG. 2 is an enlarged perspective view of the film of FIG. 1 and further comprising a functional layer applied to one surface thereof;
 FIG. 3 is a schematic diagram of a film sample used to determine tear resistance of multilayer films according to the invention in a Graves area test;
 FIG. 4 is a graphical representation of stress vs. Graves elongation in a Graves area test for three different films;
 FIG. 5 is a graphical representation of Graves area vs. weight % of a sebacic acid based copolyester for several films according to the invention;
 FIG. 6 is a graphical representation of Graves area vs. film thickness for several films according to the invention and several comparative films;
 FIG. 7 is an enlarged, vertical sectional view of a glazing unit according to the invention which includes a security control laminate bonded to a glazing member, the laminate comprising two multilayer tear resistant films according to the invention;
 FIG. 8 is an enlarged, vertical sectional view of a second embodiment of a glazing unit according to the invention and similar to FIG. 7 but further including a metalized layer;
 FIG. 9 is an enlarged, vertical sectional view of a third embodiment of a glazing unit according to the invention and similar to FIG. 7 but utilizing a single multilayer tear resistant film according to the invention; and
 FIG. 10 is an enlarged, vertical sectional view of a security control laminate similar to that illustrated in FIG. 8 but secured to a removable release liner.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The invention relates to tear resistant multilayer films comprising interdigitated layers of at least one ductile sebacic acid based copolyester (sometimes referred to herein as the "ductile" material), at least one stiff polyester or copolyester (referred to herein sometimes as the "stiff" material) and, optionally, at least one intermediate material. The exact order of the individual layers is not critical provided that at least one layer of a stiff polyester/copolyester and at least one layer of a ductile sebacic acid based copolyester are present.

Examples of some film structures within the scope of the invention include:

$S(DS)_x$
 $D(SD)_x$
 $D(ISID)_y$
 $S(IDIS)_y$

wherein S is the stiff polyester/copolyester, D is the ductile sebacic acid based copolyester, I is the optional intermediate material, x is a whole number of at least 1 (preferably at least 2 and more preferably about 6), and y is a whole number of at least 1 (preferably at least 2 and more preferably about 3). Other layer arrangements in which the order is essentially random are also possible. The two outer layers may be the same or may be different. The individual stiff polyester/copolyester layers may be comprised of the same or different materials so long as the materials are stiff. Similarly, the individual ductile sebacic acid based copolyester layers may be comprised of the same or different materials. Preferably, each stiff layer is provided by the same material and each ductile layer is the same so as to facilitate film production.

A film 10 according to the invention and having the structure $D(ISID)_y$, where y is 2 is shown in FIG. 1. Film 10 includes 9 alternating layers of a ductile sebacic acid based copolyester 11, an intermediate material 12, and a stiff polyester/copolyester 13. The two outer layers are formed of ductile sebacic acid based copolyester 11. However, the structure of FIG. 1 could be such that either stiff polyester/copolyester 13 or intermediate material 12 provides the outer layers. Preferably the film comprises at least 3 layers, more preferably from 5 to 35 layers, and most preferably about 13 layers, although as many layers as desired (e.g., 61 layers) may be employed.

The thickness of each layer and the total thickness of the film may be varied over wide limits within the scope of the invention. The practical thickness of the film is limited only by the handling characteristics desired. The lower useful practical limit is that at which the film becomes too flimsy to be readily handled or is no longer sufficiently tear resistant while the upper useful limit is that at which the film becomes overly rigid and too difficult to process. Within these constraints, films according to the invention typically have a nominal thickness in the range of from about 7 to 500 microns

(i.e., micrometers) (μm) and, more preferably, from about 15 to 185 μm .

The thickness of the individual layers may also vary over a wide range, it being understood that as the number of layers increases at a constant or decreasing film thickness, the thickness of each layer declines. The individual layers of stiff polyester/copolyester typically have an average nominal thickness of at least about 0.5 μm , more preferably from 0.5 μm to 75 μm , and most preferably from about 1 to 25 μm . Although the thickness of each layer may be the same, it is preferred that the ductile sebacic acid based copolyester layers be thinner than the stiff polyester/copolyester layers. The ductile sebacic acid based copolyester layers may range in average nominal thickness from greater than about 0.01 μm to less than about 5 μm , more preferably, from about 0.2 to 3 μm . All film and layer thickness stated herein are nominal thicknesses which may be measured according to the procedure set forth in ASTM Test Method D 1004.

Stiff polyesters/copolyesters useful in the practice of the invention comprise the reaction product of dicarboxylic acid (including ester derivatives thereof) and diol components. Preferably, the dicarboxylic acid component is either terephthalic acid or naphthalene dicarboxylic acid (such as dimethyl 2,6-naphthalene dicarboxylic acid), and the diol component is either ethylene glycol or 1,4-butanediol. Accordingly, preferred polyesters include polyethylene terephthalate, polyethylene naphthalate, polybutylene terephthalate, and polybutylene naphthalate, as well as blends thereof.

Additional stiff copolyesters based on these materials may also be used by copolymerizing the terephthalic and/or naphthalene dicarboxylic acid component(s) with one or more other diacids, including adipic, azelaic, sebacic, isophthalic, dibenzoic and cyclohexane dicarboxylic acids. Similarly, various stiff copolyesters may be formed by copolymerizing the ethylene glycol and/or 1,4-butanediol component(s) with one or more other diols such as diethylene glycol, propanediol, polyethylene glycol, polytetramethylene glycol, neopentyl glycol, cyclohexane dimethanol, 4-hydroxy diphenol, bisphenol A, and 1,8-dihydroxy biphenyl. Useful stiff polyester/copolyester materials may also be provided by incorporating one or more other diacids and/or one or more other diols into the polymerization mixture. The amount of such other materials may be varied over wide limits so long as the resulting polyester/copolyester is stiff.

As used herein, "stiff" means stretch resistant, creep resistant and dimensionally stable. More particularly, "stiff" polyesters and copolyesters according to the invention are high tensile modulus materials, preferably materials having a tensile modulus, at the temperature of interest, greater than 200 kpsi (kpsi = 1000 pounds per square inch = 6.9 MPa) (1,380 megaPascals (MPa)), more preferably greater than 300 kpsi (2,070 MPa), and most preferably greater than 400 kpsi (2,760 MPa). In some instances, orientation may be necessary to achieve the desired tensile modulus.

Tensile modulus of the stiff polyester/copolyester is determined according to ASTM Test Method D 822-88 using a 4 inch (10.2 centimeters (cm)) gauge length and a separation rate of 2 inches/minute (5 cm/min). The "temperature of interest" means the average temperature at which the film (or a structure incorporating the film) is intended to be used. ASTM D 882-88 specifies a test temperature of $23^\circ\text{C} \pm 2^\circ\text{C}$. If the temperature of interest for the multilayer film is within this range, the ASTM test procedure is followed as published. If, however, the temperature of interest is outside this range, then the test procedure is followed with the exception that the test is performed at the temperature of interest.

Ductile sebacic acid based copolyesters useful in the invention generally have a tensile modulus of less than 200 kpsi (1,380 MPa) and a tensile elongation, at the temperature of interest as defined above, of greater than 50%, preferably greater than 150%. Tensile modulus and tensile elongation of the ductile material are measured in accordance with ASTM Test Method D 882-88, a tensile test, using a 4 inch (10.2 cm) gauge length and a separation rate of 5 inches/minute (12.7 cm/min). "Tensile elongation," as used herein, refers to the elongation at break of the ductile material as measured during the referenced tensile test procedure.

Ductile sebacic acid based copolyesters according to the invention generally comprise the reaction product of terephthalic acid and/or naphthalene dicarboxylic acid such as dimethyl 2,6-naphthalene dicarboxylic acid (or ester derivatives thereof), sebacic acid (or ester derivatives thereof), and ethylene glycol. Additional sebacic acid based copolyesters may be made by polymerizing these acids with one or more other diacids such as isophthalic acid, adipic acid, azelaic acid, and cyclohexane dicarboxylic acid. Similarly, the ethylene glycol may be polymerized with one or more other diols such as diethylene glycol, propanediol, butanediol, neopentyl glycol, polyethylene glycol, polytetramethylene glycol, poly ϵ -caprolactone, polyester glycol and cyclohexane dimethanol. The relative amounts of the diacid and diol components may be varied over wide limits so long as the resulting sebacic acid based copolyester remains ductile.

The ductile sebacic acid based copolyester may comprise from 20 to 80 mole equivalents of terephthalic acid and, correspondingly, 80 to 20 mole equivalents of sebacic acid to provide the dicarboxylic acid component, and 100 mole equivalents of ethylene glycol for the diol component. (As used herein, mole equivalents and mole % are the same as the reactive systems are based on 100 equivalents.) At increasing amounts of sebacic acid, it may be more difficult to manufacture the ductile material using conventional polyester resin processing techniques. Consequently, a particularly preferred ductile sebacic acid based copolyester comprises 70 to 50 mole equivalents of terephthalic acid and, correspondingly, 30 to 50 mole equivalents of sebacic acid to provide the dicarboxylic acid component, and 100 mole equivalents of ethylene glycol for the diol component. Most preferably, the terephthalic acid provides 60 mole equivalents and the sebacic acid provides 40 mole equivalents. In another preferred ductile sebacic acid based copolyester, a portion of the sebacic acid is replaced with cyclohexane dicarboxylic acid which, with the sebacic acid, provides 20 to 80, more

preferably 30 to 50, and most preferably 40 mole equivalents.

It has been found that relatively small amounts of the ductile sebacic acid based copolyester (i.e., amounts of less than 5 weight percent), relative to the stiff polyester/copolyester, can greatly improve the tear resistance of multilayer films made therewith. However, as little as about 1 weight percent (weight % or wt. %), preferably at least about 2.6 weight %, of the ductile sebacic acid based copolyester is believed to be sufficient. Sebacic acid based copolyester material loadings up to about 10 to 20 weight % may be used although exceeding this range may reduce the tear resistance of films made therewith.

Preferably, films according to the invention have an interlayer adhesion of at least 0.1 pounds/inch width (piw) (18 grams/cm (g/cm)) more preferably at least 0.5 piw (90 g/cm). Peel adhesion may be tested using ASTM Test Method F904-84 with a separation rate of 2 inches/minute (5 cm/min.). What constitutes an acceptable interlayer adhesion will be dictated in large part by the application intended for the multilayer film. Thus, if the film provides the backing for an abrasive sanding disc, which may encounter high shear forces in use, an interlayer adhesion of at least 1 piw (180 g/cm), preferably at least 3 piw (540 g/cm), may be necessary. On the other hand, for static single use applications such as shatterproof or anti-lacerative window films, less interlayer adhesion such as 0.01 piw (2 g/cm) may be acceptable. More or less interlayer adhesion may be desirable depending on the failure mode of the film as it tears.

Because films of the invention comprise a number of interleaved layers of different materials, it is sometimes necessary to provide a means for increasing the interfacial adhesion between adjacent layers to achieve the desired interlayer adhesion. Several techniques may be used. For example, when the interfacial adhesion between adjacent layers of stiff polyester/copolyester and ductile sebacic acid based copolyester is considered inadequate, a low concentration (e.g. about 0.01 to 10%) of a component which contains an appropriate functional group may be incorporated into either or both of the ductile and stiff materials to promote interlayer adhesion. This may be accomplished by, for example, reacting or blending the functional group-containing component with the ductile or stiff material or by copolymerizing or blending it with the monomers used to provide the ductile or stiff material. Examples of useful adhesion-promoting, functional group-containing components include acrylic acid, methacrylic acid, maleic anhydride, vinyl pyridine, oxazoline-containing materials (such as polyethyl oxazoline), and the like.

Alternatively, a layer of an appropriate intermediate material may be utilized as a tie layer between the layers of stiff polyester/copolyester and ductile sebacic acid based copolyester. The intermediate layer may comprise a ductile material, a stiff material, or a rubbery material. The intermediate layer could also comprise a blend of stiff and ductile materials. Ductile and stiff are defined above. Rubbery materials manifest no significant yield point, but typically display a sigmoidal rise in elongation with applied load until rupture occurs at high strain. Whatever the precise nature of the intermediate material, if it is being used as a tie layer, it must enhance the adhesion between the stiff polyester/copolyester and ductile sebacic acid based copolyester materials. Combinations of these approaches, or even other approaches may also be used.

Many materials are useful as the intermediate layer. They include ethylene/vinyl acetate copolymers, preferably containing at least about 10% by weight vinyl acetate and a melt index of about 10, e.g., the ELVAX series of materials (duPont); carboxylated ethylene/vinyl acetate copolymers, e.g., CXA 3101 (duPont); copolymers of ethylene and methyl acrylate, e.g., POLY-ETH 2205 EMA (available from Gulf Oil and Chemicals Co.), and ethylene methacrylic acid ionomers e.g., SURYLN (duPont); ethylene/acrylic acid copolymers; and maleic anhydride modified polyolefins and copolymers of polyolefins, e.g., MODIC resins (available from Mitsubishi Chemical Company).

Other materials useful as the intermediate layer include polyolefins containing homogeneously dispersed vinyl polymers such as the VMX resins available from Mitsubishi (e.g., FN-70, an ethylene/vinyl acetate-based product having a total vinyl acetate content of 50% and JN-70, an ethylene/vinyl acetate-based product containing 23% vinyl acetate and 23% dispersed poly(methyl methacrylate)), POLYBOND (believed to be a polyolefin grafted with acrylic acid) available from Reichold Chemicals Inc., and PLEXAR (believed to be a polyolefin grafted with polar functional groups) available from Chemplex Company. Also useful are copolymers of ethylene and methacrylic acid such as the PRIMACOR family available from Dow Chemical Co. and NUCREL available from duPont. Other ethylene copolymers such as ethylene/methyl methacrylate, ethylene/ethyl acrylate, ethylene/ethyl methacrylate and ethylene/n-butyl acrylate may be used.

Various polyesters and copolyesters may also function as an intermediate layer.

The intermediate layer preferably comprises from about 1 to 30 (most preferably from about 2 to 10) weight % of the film. The nominal thickness of the intermediate layer can vary over a wide range depending on the number of layers in the multilayer film and the overall thickness of the film, but preferably is from about 0.01 μm to less than about 5 μm , more preferably from about 0.2 to 3 μm .

Alternatively, adjacent layers of the stiff and ductile materials may be treated with radiation, such as ultraviolet, electron beam, infrared or microwave radiation, to improve interlayer adhesion.

Each of the stiff, ductile and intermediate layer materials may further include or be supplemented with various adjuvants, additives, colorants, extenders, antioxidants, thermal stabilizers, ultraviolet light stabilizers, plasticizers, slip agents, etc. that are conventionally and customarily used in the manufacture of such materials or films made therewith.

These supplemental materials may comprise up to about 5 weight % of the total weight of the layers into which they are incorporated so long as the tear resistance of the film is not significantly adversely affected.

If desired, a functional layer may be applied to one or both of the major surfaces of the film. For example, an adhesive 14 may be applied to at least one of the major surfaces as shown in FIG. 2. Adhesive 14 may be activatable by pressure, heat, solvent or any combination thereof and may be of any type such as an acrylate, a rubber/resin, or a silicone. Other functional layers, for example, an abrasive material (optionally in a binder), a radiation (e.g., light) sensitive or blocking layer, an ink-receptive layer, a magnetic recording media, a top coat, a slip agent layer, a vapor coated material, a primer layer, a reflective layer, or a moisture or gas barrier layer may be employed. Other functional layers may also be used. The functional layers may be employed singly or in combination with other functional layers on one or both sides of the film.

To modify the surface properties of the film or to promote adhesion of any subsequently applied functional layer, the film may be pretreated with a primer coating, activated by flame or corona discharge or other surface treatments, or a combination of these approaches.

Films according to the invention may be readily made using techniques known in the art. One such technique is disclosed in U.S. Patent No. 3,565,985 (Schrenk et al.). In making films of the invention, melt coextrusion by either the multimanifold die or the feedblock method in which individual layers meet under laminar flow conditions to provide an integral multilayer film may be used. More specifically, separate streams of the ductile, stiff and, optionally, intermediate materials in a flowable state are each split into a predetermined number of smaller or sub-streams. These smaller streams are then combined in a predetermined pattern of layers of stiff, ductile and, optionally, intermediate materials to form an array of layers of these materials in a flowable state. The layers are in intimate contact with adjacent layers in the array. This array generally comprises a tall stack of layers which is then compressed to reduce its height. In the multimanifold die approach, the film width remains constant during compression of the stack while the width is expanded in the feedblock approach. In either case, a comparatively thin, wide film results. Layer multipliers in which the resulting film is split into a plurality of individual subfilms which are then stacked one upon another to increase the number of layers in the ultimate film may also be used.

In manufacturing the films, the materials may be fed such that any one of the three constitutes the outer layer. The two outer layers often comprise the same material. Preferably, the materials comprising the various layers are processable at the same temperature and have similar melt viscosities so as to avoid degrading a lower melting material. Accordingly, residence time and processing temperatures may have to be adjusted depending on the characteristics of the materials of each layer.

Other manufacturing techniques such as lamination, coating or extrusion coating may be used in assembling multilayer films according to the invention. For example, in lamination, the various layers of the film are brought together under temperature and/or pressure (e.g., using heated laminating rollers or a heated press) to adhere adjacent layers to each other. In extrusion coating, a first layer is extruded onto either a cast web, a monoaxially oriented film or a biaxially oriented film and subsequent layers are sequentially coated onto the previously provided layers. Exemplary of this method is U.S. Patent No. 3,741,253. Extrusion coating may be preferred over the melt coextrusion process described above where it is desirable to pretreat selected layers of the multilayer film or where the materials are not readily coextrudable.

It is preferred that the polyester/copolyester layers be oriented, either uniaxially or biaxially, at a temperature above their glass transition temperature so as to enhance the stiffness, modulus and creep resistance of the film. (For some uses, such as thermoforming applications, orientation of the stiff polyester/copolyester layers would not be required.) Orientation of the ductile sebacic acid based copolyester and intermediate layer materials is optional. Orientation may be accomplished by conventional methods typically used in the art such as mechanical stretching (drawing) or tubular expansion with heated air or gas. Typical draw ratios are in the range of 2.5 to 6 times in either or both of the machine and transverse directions. Greater draw ratios (for example, up to about 8 times) may be used if the film is oriented in only one direction. The film need not be stretched equally in the machine and transverse directions although this is preferred if balanced properties are desired.

The films may also be heat set by exposing the film to a temperature of about 10° to 150°C below the melting temperature of the stiff component for about 4 to 15 seconds so as to increase the crystallinity, stiffness, modulus and creep resistance of the film while reducing its tendency to shrink. In applications where film shrinkage is not of significant concern, the film may be heat set at relatively low temperatures or not at all. On the other hand, as the temperature at which the film is heat set is increased, the tear resistance of the film may change. Thus, the actual heat set temperature and time will vary depending on the composition of the film and perhaps its intended application but should not be selected so as to substantially degrade the tear resistant properties of the film. Within these constraints, a heat set temperature of about 135° to 205°C is generally desirable for many of the applications in which the multilayer films of the invention are useful.

Various functional layers may be subsequently applied by lamination, extrusion coating or other known techniques. Various primers and/or surface treatments may be required as discussed more fully above.

Multilayer films according to the invention are both stiff (dimensionally stable, high modulus) and tear resistant. As explained above, stiff, high tensile modulus, creep resistant films such as cellophane, polyester and biaxially oriented polypropylene packaging films have little tear resistance. On the other hand, low tensile modulus, ductile materials such as polyolefin trash bags are tear resistant but are not dimensionally stable (i.e., they stretch readily). Films according to the invention provide the desirable properties of both high tensile modulus, stiff, dimensionally stable, creep resistant materials and low tensile modulus, ductile, tear resistant materials in a multilayer arrangement. As a result, multilayer films according to the invention offer both excellent tear resistance and dimensional stability. This beneficial amalgamation of properties is achieved because the different materials which comprise the films of the invention are assembled in a multilayer arrangement. As exemplified below, single layer blends of stiff polyester/copolyester materials and ductile sebacic acid based copolyester materials do not equally reflect the characteristics of films according to the invention.

The tear resistance of films according to the invention may be measured by ASTM Test Method D 1004 (also known as a Graves tear test). In a Graves tear test, a film sample 16 having the general shape shown in FIG. 3 (and described more explicitly in ASTM D 1004) is clamped between opposed jaws with an initial separation of 1 inch (2.5 cm). The jaws are then moved apart at a constant rate of 2 inches/minute (5 cm/min.) to tear the film in the area of the sample designated by the reference numeral 18. The tearing stresses imposed on the film are concentrated in area 18. The film may be torn in either the machine direction (i.e., the direction in which the film is extruded) or the transverse direction (i.e., perpendicular to the machine or extrusion direction). The test direction corresponds to the orientation of area 18. More specifically, a pair of axes labeled A-B and C-D have been superimposed on film sample 16 in FIG. 3. The opposed jaws are moved along axis A-B to tear film sample 16 along axis C-D.

With reference to FIG. 4, test data were recorded by graphically plotting the stress (as measured in kpsi) experienced by the film versus the strain (as measured by Graves elongation in %) that the film underwent during the test. "Stress" is defined as the recorded force divided by the product of the film thickness and the ligament width (distance "d" in FIG. 3). The expression "Graves elongation" as used herein refers to the elongation of a film in the tear direction as observed during a Graves area test and reflects the percent change in the jaw separation distance that occurs during the test relative to the jaw separation distance at the outset of the test. "Graves elongation at break" as used herein refers to the elongation of the film in % at its break point observed during the Graves area test. (It will be understood that Graves elongation at break differs from tensile elongation. Tensile elongation is measured during a tensile test and may be used to characterize ductile materials useful in the invention as explained above.)

With continued reference to FIG. 4, the plot (i.e., "curve") labeled with the letter "A" describes a film having a large maximum stress which falls off quickly as the film is stretched during the test. Curve A typifies the performance of a high modulus, stiff, dimensionally stable material which has poor tear resistance (as shown by the rapid falloff in stress as the film tears). Polyesters, cellophane, biaxially oriented polypropylene and similar packaging films perform similarly to curve A. Curve A was obtained by measuring the performance of the polyethylene terephthalate film of comparative example 11, described more fully below.

The curve labeled with the letter "B" describes the performance of a low modulus, ductile, readily stretchable, traditionally tear resistant material (as evidenced by the relatively high Graves elongation at break relative to curve A) because the film stretches rather than tears. The film is capable of sustaining only a relatively low stress. Plastic trash and grocery bags are common examples of films that would perform in a manner similar to that described by curve B. Curve B was obtained by measuring the performance of the linear low density polyethylene film of comparative example 12, described more fully below.

Curve "C" illustrates the performance of a multilayer film according to the invention and, more specifically, the film of example 27 described below. The maximum stress sustained by this film is similar to or exceeds the stiff film of curve A. However, the stress experienced by the curve C film of the invention does not fall off as rapidly as in the case of the curve A film. Thus, as compared to conventional polyester films of curve A, films according to the invention are more able to successfully withstand catastrophic tearing forces while being of substantially equal modulus. Such a property is highly desirable in certain applications, especially shatterproofing film for windows where the impact from breaking glass may be sudden and catastrophic. As compared to the low modulus films of curve B, films of the invention are able to sustain much higher stress. Thus, films according to the invention are both stiff (high modulus) and tear resistant.

In a Graves tear test, tear resistance data are conventionally reported as the maximum force experienced by the film. The data reported herein, however, are the total area (referred to herein at times as the "Graves area") beneath the stress-strain curve (i.e., the curves of FIG. 4) which is obtained by a mathematical integration of the curve. Graves area is regarded as a measure of the total energy required to cause the film to fail and, hence, a measure of the film's combined stiffness and tear resistance. Thus, Graves area may be regarded as a measure of the ability of the film to absorb energy. Graves area is reported herein in units of kpsi% wherein 1 kpsi% = 69 kilojoules/cubic meter. It will be understood that films with a relatively large Graves area have enhanced combined stiffness and tear resistance relative to those films with a relatively small Graves area.

As shown more fully below, Graves area may vary depending on whether the test is conducted in the machine or the transverse direction of the film. Also, Graves area generally increases with increasing film thickness. As a general

characterization, a multilayer film may be regarded as tear resistant within the scope of the invention if it demonstrates a Graves area in one direction which exceeds the Graves area (in the same direction) of a single layer film that comprises only the stiff polyester or copolyester used in the multilayer film, the single layer film being processed (i.e., oriented, heat set, etc.) in the same manner as the multilayer film and to a substantially equal film thickness. Preferably and more specifically, a multilayer film may be regarded as tear resistant within the scope of the invention if it demonstrates a Graves area at least equal to $40 + 0.4(x)$ kpsi% in one direction (e.g., the machine or the transverse direction) of the film, wherein x is the nominal thickness of the film in microns.

Furthermore, and related to their overall performance, multilayer films of the invention preferably have a tensile modulus (when tested according to ASTM Test Method D 882-88) of at least 175 kpsi (1,208 MPa) in one direction of the film, more preferably at least 240 kpsi (1,656 MPa), and most preferably at least 450 kpsi (3,105 MPa). However, the actual modulus which is desirable will depend on the application for which the film is intended, some applications preferring relatively stiffer films and others preferring relatively more flexible films. In addition, and also related to their overall performance, multilayer films according to the invention desirably demonstrate a Graves elongation at break of at least 20%, more preferably at least 40% in the tear direction of the film measured during the Graves area test.

The invention will be more fully appreciated with reference to the following, non-limiting examples.

Examples 1 to 26

A series of multilayer films comprising alternating layers of a stiff polyester/copolyester and a ductile sebacic acid based copolyester was formed. A stiff polyethylene terephthalate (PET) (differential scanning calorimetry (DSC) melting point of 256°C; intrinsic viscosity of 0.60 deciliters per gram (dl/g) as measured in 60% phenol and 40% dichlorobenzene at 110°C) was coextruded with a ductile sebacic acid based copolyester. The sebacic acid based copolyester comprised 40 mole % sebacic acid and 60 mole % terephthalic acid as the dicarboxylic acid components and 100 mole % ethylene glycol as the diol component. The sebacic acid based copolyester had an intrinsic viscosity in the range of 0.9 to 1.05 dl/g when measured in the same fashion as the PET. The ductile sebacic acid based copolyester also displayed a tensile modulus of 14 kpsi (97 MPa) and a tensile elongation of 355% when tested according to ASTM D822-88 at room temperature using the ductile material test protocol described above.

The multilayer films were coextruded onto a chilled casting wheel and subsequently oriented sequentially 2.6 times in the machine direction (MD) at 80°C and 4.2 times in the transverse direction (TD) at 99°C. The films were then heat set at 149°C.

The number of layers, the film thickness, and the weight percent of the ductile sebacic acid based copolyester were varied as shown below in Table 1. The tear resistance of the films in both the machine and the transverse directions are reported below in Table 1 as Graves area (rounded to the nearest 10 here and for other examples) according to the procedure described more fully hereinabove. The Graves elongation at break values (rounded to the nearest 5 here and for other examples) are also reported in Table 1. The reported Graves area and Graves elongation at break values throughout the application (unless noted otherwise) are an average of 9 readings in each of the machine and transverse directions.

Although ASTM D 1004 utilizes a 0.5 inch (1.3 cm) ligament (distance "d" in FIG. 3), examples 1 to 26 herein were analyzed using a 1.31 inch (3.3 cm) ligament. For examples 1 to 26, the observed results were mathematically converted to a value corresponding to a 0.5 inch (1.3 cm) ligament by multiplying the observed Graves area result by 0.678 and adding 32.4, this conversion factor having been determined by a linear regression analysis of multiple samples. The observed results for Graves elongation at break for examples 1 to 26 were also mathematically converted so as to correspond to a 0.5 inch (1.3 cm) ligament by multiplying the observed result by 0.655 and adding 11.3, this conversion factor having been determined by a linear regression analysis of multiple samples.

Table 1

Example	Number of Layers	Film Thickness (μm)	Wt. % of Ductile Material	Graves Area (kpsi%)		Graves Elongation at Break (%)	
				MD	TD	MD	TD
1	5	54.6	10	160	160	40	40
2	5	54.1	20	110	130	30	35
3	5	47.8	30	120	130	35	40
4	5	45.7	40	80	110	25	40
5	5	45.7	50	60	80	20	35

Table 1 (continued)

Example	Number of Layers	Film Thickness (μm)	Wt. % of Ductile Material	Graves Area (kpsi%)		Graves Elongation at Break (%)	
				MD	TD	MD	TD
6	13	45.7	40	190	190	30	40
7	13	57.7	40	240	210	45	45
8	13	54.6	20	160	150	30	40
9	13	49.5	30	80	100	25	35
10	13	53.5	40	90	90	30	35
11	13	48.3	50	70	80	25	30
12	29	45.7	10	190	130	30	30
13	29	46.5	20	110	80	30	30
14	61	47.5	10	130	100	30	30
15	61	53.3	20	120	80	30	25
16	61	53.3	30	100	80	25	30
17	61	53.3	40	80	70	25	25
18	61	50.8	50	70	80	25	25
19	5	27.9	40	130	120	25	30
20	13	25.4	10	140	130	30	30
21	29	26.2	10	90	70	20	20
22	61	24.1	10	110	80	25	20
23	5	15.2	10	90	90	20	25
24	13	14.0	10	80	70	20	20
25	29	15.2	10	80	70	20	20
26	61	12.7	10	60	50	15	15

The data of Table 1 show that as the number of layers in the film remains constant, the Graves area of the film decreases as the amount of ductile sebacic acid based copolyester increases above 10%. The data of Table 1 further show that as the total number of layers increases, the tear resistance of the films tends to increase and then becomes more constant or decreases as the number of layers approaches 61, especially at lower wt. % amounts of the ductile sebacic acid based copolyester. Consequently, films according to the invention comprise at least 3 layers, more preferably from 5 to 35 layers, and most preferably about 13 layers.

FIG. 5 is a graphical representation of the data of examples 1 to 18, the plotted Graves area being an average of the MD and TD values from Table 1. FIG. 5 illustrates the relationship among Graves area, the wt. % of the ductile sebacic acid based copolyester, and the number of layers in the film as the total film thickness was attempted to be held relatively constant. Using linear regression analysis, the lines which "best fit" the data sets (based on the number of layers in the film) were drawn.

However, as shown in examples 19 to 26, tear resistance is also related to film thickness and the above trends may not always be rigidly observed as film thickness decreases. Thicker multilayer films generally have enhanced tear resistance relative to thinner multilayer films when the number of layers and the amount of ductile material are essentially constant.

Comparative Examples 1 to 6

Comparative examples (C.E.) 1 to 6 report a series of single layer films formed by extruding the PET of examples 1 to 26 onto a chilled wheel. The films were sequentially drawn 3.5 to 4 times in the machine direction at about 85 to 90°C, and then about 4.5 times in the transverse direction at about 100°C. The films were subsequently heat set at 220

to 225°C. The films so produced were regarded as representative of conventional, commercially available PET films such as might be used in packaging applications. The films were tested for Graves area in both the machine and transverse directions according to the procedure described above and with the results shown below in Table 2.

(The processing conditions in the preparation of these comparative examples were not identical to those used in preparing examples 1 to 26. It will be understood by those of ordinary skill in the art that adjustments in processing conditions can affect film properties. However, the films of comparative examples 1 to 6 are regarded as representative of conventional, commercially available PET films. Other comparative data which replicate examples herein may be found. See example 27, for instance.)

TABLE 2

Example	Film Thickness (μm)	Graves Area (kpsi%)		Graves Elongation at Break (%)	
		MD	TD	MD	TD
C.E. 1	11.9	40	30	5	10
C.E. 2	22.4	60	40	10	10
C.E. 3	35.0	40	50	10	15
C.E. 4	45.5	40	50	15	10
C.E. 5	96.3	70	70	15	20
C.E. 6	174.0	90	80	20	20

The data of examples 1, 6, 7, 12, 14, and 19 to 26 were graphically plotted in FIG. 6 to illustrate the relationship among Graves area, film thickness, and the number of layers as the wt. % of the ductile material was held constant at 10%. Separate curves were then constructed for the 5, 13, 29 and 61 layer films in both the machine and transverse directions by serially connecting the data points. Separate curves were also prepared in the machine and transverse directions for the single layer PET films of comparative examples 1 to 5. (Comparative example 6 was not included in FIG. 6 in order to facilitate data management and presentation of the graph.) As shown in FIG. 6, multilayer films according to the invention, virtually without exception, demonstrated a Graves area which exceeded that observed for the conventional PET films of comparative examples 1 to 5, whether tested in the machine or the transverse direction.

Also shown in FIG. 6 is the line defined by the equation $40 + 0.4(x)$ kpsi% wherein x is the nominal thickness of the film in microns. Multilayer films according to the invention have Graves area values which generally fall above this line whereas the conventional PET films of comparative examples 1 to 5 have Graves area values which generally fall below this line. Thus preferred multilayer films comprising alternating layers of a stiff polyester/copolyester, a ductile sebacic acid copolyester and, optionally, an intermediate material, according to the invention are considered to be tear resistant if they demonstrate a Graves area which is equal to or which exceeds $40 + 0.4(x)$ kpsi% wherein x is the nominal thickness of the film in microns. As explained above and related to their overall performance, tear resistant films of the invention also preferably exhibit a tensile modulus (as determined in a conventional tensile test) in one direction of the film of at least 175 kpsi (1,203 MPa), more preferably at least 240 kpsi (1,650 MPa), and most preferably at least 450 kpsi (3,105 MPa), as well as a Graves elongation at break (as determined during the assessment of Graves area) of at least 20%, preferably at least 40%.

Example 27

A film comprising a total of 13 alternating layers of the stiff PET of examples 1 to 26 coextruded with 5 weight % of the ductile material of the same examples was prepared. The film was cast onto a chilled quenching wheel, sequentially oriented 2.6 times in the machine direction at 86°C and 4.5 times in the transverse direction at 103°C, and heat set at 149°C. The film was about 62 μm thick and displayed a Graves area of 330 kpsi% in the machine direction and 220 kpsi% in the transverse direction. The film also exhibited a tensile modulus of 500 kpsi in the machine direction and 700 kpsi in the transverse direction. The Graves elongation at break was 45% in each of the machine and transverse directions. The multilayer film of this example was used to prepare curve C of FIG. 4.

When a single layer film comprising only the stiff PET of this example was extruded, biaxially drawn and heat set in the same manner at a thickness of about 66 μm , it demonstrated a Graves area of 120 kpsi% in the machine direction and 80 kpsi% in the transverse direction. The single layer film also displayed a tensile modulus of 530 kpsi in the machine direction and 730 in the transverse direction and a Graves elongation at break of 30% in each direction. (Data

reported for the single layer PET film are an average of 5 measurements in each direction.) Even though the tensile moduli of the multilayer and single layer films were comparable, the multilayer films demonstrated superior tear resistance as measured by the Graves area test.

5 Comparative Examples 7 to 10

A series of comparative examples was prepared by extruding the stiff and ductile materials of examples 1 to 26 into a blended single layer film rather than a multilayer film. The single layer films were extruded onto a chilled casting wheel, biaxially oriented 3.3 times in each of the machine and transverse directions at 100°C, and heat set at 140°C. The weight % of the ductile material was varied as shown below in Table 3 along with the results of the Graves area and Graves elongation at break tests, the reported data being an average of 5 measurements in each direction. The films were not sufficiently uniformly thick to permit Graves area, tensile modulus, and Graves elongation at break testing at one thickness in both the machine and transverse directions. Consequently, Table 3 also reports the film thickness for testing in each direction, the reported thicknesses being an average of 5 measurements in each direction.

Table 3

Example	Wt. % of Ductile Material	Film Thickness (μm)		Tensile Modulus(kpsi)		Graves Area (kpsi%)		Graves Elongation at Break(%)	
		MD	TD	MD	TD	MD	TD	MD	TD
C.E. 7	0	33	39	640	640	70	100	20	30
C.E. 8	5	43	25	420	450	80	110	30	35
C.E. 9	10	27	38	440	570	80	80	30	25
C.E. 10	30	80	59	410	420	80	110	30	35

Comparative examples 7 to 10 illustrate that blends of stiff and ductile materials extruded as single layer films do not exhibit any significant improvement in tear resistance with the addition of a ductile sebacic acid based copolyester. This is in distinction to the benefits which are achieved by coextruding the stiff and ductile materials into a multilayer film according to the invention.

35 Comparative Example 11

Comparative example 11 describes the preparation of the single layer PET film measured by curve A of FIG. 4. More specifically, the PET of examples 1 to 26 was melt extruded onto a chilled casting wheel and then sequentially oriented 3.4 times in the machine direction at 88°C and 4.0 times in the transverse direction at 110°C, followed by heat setting at 232°C. The finished film was 51 μm thick and demonstrated a Graves area of 30 kpsi% in the machine direction and 37 kpsi% in the transverse direction as well as a tensile modulus of 660 kpsi in the machine direction and 650 kpsi in the transverse direction. The film of this example is considered representative of a conventional biaxially oriented PET film.

45 Comparative Example 12

Comparative example 12 describes the preparation of the single layer linear low density polyethylene film measured by curve B of FIG. 4. More specifically, TF0119F linear low density polyethylene (hexene comonomer) having a density of 0.918 grams/cubic centimeter and commercially available from Novacor Chemicals, Inc. (Calgary, Alberta) was extruded and blown into a 51 μm thick film. The blow up ratio was 3.2 and the draw down ratio was 12.3. The film demonstrated a Graves area of 182 kpsi% in the machine direction and 200 kpsi% in the transverse direction due significantly to the large Graves elongation (greater than 180%). However, the film exhibited a relatively low stress. The film of this example is considered representative of films conventionally employed in the manufacture of garbage and grocery bags.

Examples 28 to 31

Comparative Examples 13 and 14

A series of examples was prepared to illustrate the improvement in tear resistance that is possible when multilayer films comprising alternating layers of stiff and ductile materials are oriented in only one direction. More specifically, a series of 13 layer films having the composition of the film of example 27 (the PET of examples 1 to 26 with 5 wt. % of the ductile sebacic acid based copolyester of the same examples) was extruded onto a chilled casting wheel. A square sample of each film was clamped on all four sides and drawn at 100°C 4.0 times in one direction at a constant width while being restrained in the transverse direction. The film was then heat set at 150°C. The tear resistance and Graves elongation at break of the film in the direction of orientation (MD) and the direction perpendicular thereto (TD) were tested as described above with the results shown below in Table 4. Also evaluated and reported in Table 4 as comparative examples 13 and 14 are two single layer films comprising the PET of example 27 processed as described for examples 28 to 31.

Table 4

Example	Film Thickness (μm)	Graves Area (kpsi%)		Graves Elongation at Break (%)	
		MD	TD	MD	TD
28	89	80	NT	30	NT
29	117	40	540	50	105
30	131	20	540	10	110
31	252	60	440	25	110
C.E. 13	90	10	NT	5	NT
C.E. 14	120	10	400	10	90
NT = Not tested					

These examples show that uniaxially oriented multilayer films according to the invention can offer improved tear resistance relative to single layer films comprising only a stiff PET.

Examples 32 to 38

A series of examples was prepared to illustrate the improvement in tear resistance that is possible when the multilayer films comprise three alternating layers of stiff and ductile materials. More specifically, the stiff PET and the ductile sebacic acid based copolyester of examples 1 to 26 were coextruded onto a chilled casting wheel in a three layer configuration in which the two outer layers comprised the stiff PET. The extruded films were then sequentially biaxially oriented 2.6 times in the machine direction at 100°C and 4.2 times in the transverse direction at 100°C. The films were then heat set at 149°C. The thickness of the films and the weight percent of the ductile sebacic acid based copolyester were varied as shown below in Table 5 along with the results from the Graves area and Graves elongation at break testing of these examples. The reported film thicknesses are an average of the thicknesses of the samples tested in the machine and transverse directions. The reported Graves area and Graves elongation at break data are an average of 5 readings in each of the machine and transverse directions.

Table 5

Example	Film Thickness (μm)	Wt. % of Ductile Material	Graves Area (kpsi%)		Graves Elongation at Break (%)	
			MD	TD	MD	TD
32	48.9	3	330	150	65	50
33	48.9	5	250	110	55	30

Table 5 (continued)

Example	Film Thickness (μm)	Wt. % of Ductile Material	Graves Area (kpsi%)		Graves Elongation at Break (%)	
			MD	TD	MD	TD
34	52.2	10	260	60	55	20
35	65.2	20	270	70	70	35
36	63.8	30	140	70	40	30
37	55.0	40	110	30	40	20
38	55.9	50	90	40	45	25

A 68.8 μm thick single layer film comprising only the stiff PET of these examples was processed under the same conditions and found to have a Graves area of 160 kpsi% in the machine direction and 110 kpsi% in the transverse direction as well as a Graves elongation at break of 40% in the machine direction and 45% in the transverse direction. Examples 32 to 38 illustrate that significant improvements in tear resistance are possible over single layer PET films when the multilayer film comprises 3 layers. Examples 32 to 38 further show the benefit of using less than 20 wt. % of the ductile sebacic acid based copolyester and the surprising improvement which is possible even when amounts of 5 wt. % or less are employed.

Example 39

A multilayer film was prepared as described in example 6 except that the ductile material comprised 60 mole equivalents terephthalic acid, 30 mole equivalents sebacic acid and 10 mole equivalents cyclohexane dicarboxylic acid. The film was coextruded onto a chilled quenching wheel, and simultaneously oriented about 3.3 times in the machine direction and about 3.5 times in the transverse direction, each at about 97°C. The film was then heat set at about 149°C. The resulting 50 μm thick film demonstrated a Graves area of 70 kpsi% in the machine direction and 60 kpsi% in the transverse direction with a Graves elongation at break of 25% in each of the machine and transverse directions.

Examples 40 to 42

Examples 40 to 42 were made in a manner similar to example 39 but comprised different mole equivalents of the sebacic acid and cyclohexane dicarboxylic acid (CHDCA) components as shown below in Table 6. The weight % of the ductile sebacic acid based copolyester was also varied as reported in Table 6 along with the results of the Graves area and Graves elongation at break testing.

Table 6

Example	Sebacic Acid Equivalents	CHDCA Equivalents	Weight % Ductile Material	Graves Area (kpsi%)		Graves Elongation at Break (%)	
				MD	TD	MD	TD
40	20	20	5	90	70	25	25
41	25	15	10	80	80	25	25
42	35	5	10	70	70	25	25

Examples 39 to 42 illustrate that a portion of the sebacic acid component may be replaced by cyclohexane dicarboxylic acid. Sebacic acid based copolyesters according to the invention may comprise, in addition to terephthalic acid, 1 to 39 mole equivalents sebacic acid and, correspondingly, 39 to 1 mole equivalents cyclohexane dicarboxylic acid, wherein the mole equivalent contributions of these three materials sums to 100.

As noted above, the combination of tear resistance and high modulus provides the multilayer films of the present invention with a unique ability to absorb energy, especially in the event of a catastrophic impact. Consequently, the multilayer films disclosed herein are useful as security control laminates for shatter-proofing glazing members against

impact or explosion. In such applications, one or more tear resistant multilayer films are applied to a glazing member as a shield that prevents the fragmentation of the glazing member even though it splinters or shatters upon breaking. When adhesively bonded to a glazing member, security control laminates based on the multilayer films of the present invention provide excellent energy absorption and distribution properties without significantly delaminating from the glazing member. The security control laminates are also less likely to puncture and/or tear.

Turning now to FIG. 7, a glazing unit 20 comprises a security control laminate 21 bonded to the interior face of a glazing member 22 by means of an adhesive layer 23, such as those commonly used for solar control or security films, including acrylate pressure-sensitive adhesives and water activated adhesives. Security control laminate 21 comprises a first multilayer film 25 (having a first face 25a and an opposed second face 25b) and a second multilayer film 27 (having a first face 27a and an opposite second face 27b) the two films being secured or bonded together by a layer of a (polyester) laminating adhesive 26.

In order to minimize the deteriorative effects of ultraviolet (UV) radiation on any of the polymeric materials which comprise the security control laminate, it is highly desirable to interpose a coating 24, containing a UV absorber, between multilayer film face 25b and adhesive layer 23. Alternatively, means for absorbing UV radiation may be incorporated into adhesive layer 23 or multilayer film 25. Suitable UV absorbent coatings may include substituted benzophenones and substituted benzotriazenes.

Multilayer film face 27b optionally includes a thin, abrasion resistant coating 28 thereon to protect film 27 from mechanical abrasion such as might occur during installation or cleaning of the security control laminate. Suitable abrasion resistant coatings comprise photopolymerized materials such as the "hydantoin hexacrylate" coatings described in U.S. Pat. No. 4,249,011 (Wendling), which is incorporated herein by reference, or other photopolymerizable multifunctional acrylates.

Although FIG. 7 illustrates the security control laminate on the interior face of the glazing member (i.e., the face of the member which is opposite to the face first exposed to the force of the impact), the laminate may also be secured to the exterior face. Also contemplated is a glazing unit comprising multiple glazing members arranged in, for example, a sandwich or an insulated construction wherein the security control laminate is secured to a face of a glazing member which is interior to the sandwich or insulated construction. Additionally, it is contemplated that the security control laminate may be adhesively or mechanically attached to a supplemental frame or batten system that surrounds the glazing member as well as to the glazing member itself. An installation of this type provides additional security against unintended removal or dislodgement of the glazing member from its frame which would otherwise allow access through the glazing unit.

FIG. 8 illustrates a second embodiment of security control laminate 21 which additionally includes a reflective metalized layer 30 to impart energy control properties to the glazing unit. This embodiment is similar to the security control laminate illustrated in FIG. 7 except that metalized layer 30 is adjacent to multilayer film face 27b rather than abrasion resistant coating 28.

More particularly, a carrier film 31 supports metalized layer 30, the latter being bonded to multilayer film face 27b by an adhesive layer 29. Metalized layer 30 may be aluminum, gold, silver, copper, nickel, or any other suitable reflector of radiant energy over the solar and infrared spectrum (i.e., a wavelength of 0.3 to 40 μm). Metalized layer 30 may be applied to carrier film 31 by, for example, vapor deposition. Preferably, metalized layer 30 is relatively transparent to visible light and offers good reflectivity of infrared radiation. Carrier film 31 comprises an optically clear film, preferably an optically clear polyester film, having a thickness in the range of about 13 to 51 μm (0.5 to 2.0 mils). Optionally, carrier film 31 may be dyed to provide additional protection from radiation incident on the glazing unit. In this regard, any optical grade dyed film may be used. Such films typically comprise an optically clear polyester film which has been dipped in a heated solvent bath containing a dye of the desired (and often customized) color, washed, rinsed and dried. Films of this type are commercially available from Martin Processing Company (Martinsville, VA). Adhesive layer 29 may be a laminating adhesive similar to adhesive layer 26 for example. The embodiment of FIG. 8 may (but need not) include abrasion resistant coating 28 on the face of carrier film 31 which does not support metalized layer 30.

FIG. 9 illustrates a third embodiment of security control laminate 21 which utilizes a single multilayer film 25. The embodiment of FIG. 9 is similar to those described in conjunction with FIGS. 7 and 8 with the exception that multilayer film 27 has been replaced by a dyed film 32 that is secured to multilayer film face 25a by way of adhesive layer 26. Abrasion resistant coating 28 protects the opposite face of dyed film 32 from mechanical abrasion. Dyed film 32 is similar to the dyed version of carrier film 31 described in conjunction with FIG. 8.

Constructions other than those illustrated above comprising different arrangements of multilayer films, metalized films, and/or dyed films are also possible. Such other constructions may offer various security and/or solar control properties as will be appreciated by the skilled artisan. Furthermore, while certain coatings and layers may be described herein as being "on" other coatings and layers of the security control laminate, it will be understood that this encompasses both direct and indirect attachment to the other coatings or layers.

FIG. 10 illustrates the security control laminate of FIG. 8 before installation on a glazing member. Overcoat 33, which preferably is a water soluble material, is applied over adhesive layer 23 to protect it from damage during manu-

facture and handling. A variety of water soluble materials such as methyl cellulose or polyvinyl alcohol are suitable as the overcoat material. Security control laminate 21 is temporarily disposed on a removable release liner 34 which is discarded prior to installation of the laminate on the glazing member. Release liners typically employed with solar control and security films may be used. The security control laminate is prepared for application by removing the release liner and rinsing the laminate with water to remove overcoat 33, thereby exposing and/or activating adhesive layer 23. Security control laminate 21 is then applied to the glazing member using conventional installation techniques known in the art.

The following examples illustrate the particular utility of multilayer films according to the invention in providing security control laminates for glazing members. In these examples all parts and percentages are by weight and all film and layer thicknesses are nominal thicknesses. Single pane window glass panels having a security control laminate according to the invention applied to one face thereof were tested for their ability to withstand impact without puncture and/or tearing in accordance with a modified version of American National Standards Institute's Specification for Safety Glazing Material Used in Buildings, ANSI Z97.1-1984. In general, tests were conducted by swinging a weighted shotbag into 0.3 cm and/or 0.6 cm thick glass panels in a pendulum arc, dropping the bag from heights of 45.7 cm (18 inches) and 122 cm (48 inches). (The shotbag impacted the surface of the glass panel which did not have the security control laminate bonded thereto.) The heights used will be recognized as corresponding to levels II and III of ANSI Z97.1-1984. The ANSI test setup was utilized but different numbers of panels were tested at less than all of the levels specified in the published procedure. An individual panel was considered to have met the test requirements, if, after impact, a 7.6 cm diameter metal ball mounted on a rod could not be passed through any break in the panel resulting from the impact.

Example 43

A multilayer film comprising 13 alternating layers of the stiff PET of examples 1 to 26 and the ductile copolyester of the same examples was coextruded onto a chilled casting wheel and subsequently sequentially oriented 2.6 times in the machine direction at about 85°C to 90°C and 3.3 times in the transverse direction at 99°C. The resulting 58 μ m thick film was heat set at 149°C and comprised 7 wt. % of the ductile material.

One surface of the multilayer film was corona treated to a surface energy of 40 to 44 dynes/cm under standard corona treating conditions using an apparatus available from Enercon Industries.

Adhesive was applied to the corona treated surface of the multilayer film by coating a 14.5% solids solution of a pressure sensitive adhesive comprising 100 parts of a 96:4 isooctyl acrylate:acrylamide copolymer (prepared as described in U.S. Pat. No. Re. 24,906 to Ulrich, which is incorporated herein by reference), 2 parts of a UV absorber (UNIVUL D-50, commercially available from BASF), 0.8 part of a fluorochemical surfactant (FC-740, commercially available from Minnesota Mining and Manufacturing Company), and 0.5 part of a substituted phenolic thioether antioxidant (SANTINOX-R, commercially available from Monsanto Company) in a solvent system comprising 33 parts heptane, 32 parts ethyl acetate, 29.5 parts toluene, and 6 parts methyl ethyl ketone. The coated multilayer film was then dried in a circulating air oven operating at 105°C for approximately 3 minutes to remove the solvent and to provide a pressure sensitive adhesive having a dry coating weight of 22.6 grams/square meter (g/m^2). A 1.6% solids overcoat solution of METHOCEL A15LV (commercially available from Dow Chemical Co.) was applied over the pressure-sensitive adhesive and the water was evaporated by passing the coated film through a circulating air oven operating at 63°C for approximately 1 minute to provide a tack-free, water soluble coating of 0.3 g/m^2 dry weight.

A 25 μ m thick release liner comprising a release agent coated PET film was removably laminated to the METHOCEL coating by passing the construction through a pair of squeeze rolls to provide a security control laminate according to the invention.

The release liner was removed, the film laminate was rinsed with water to remove the METHOCEL coating, and the security control laminate was applied to both 0.3 cm thick and 0.6 cm thick clean glass panels measuring 86.4 cm by 193 cm using standard installation techniques for solar and security films. The resulting panels were dried at room temperature for six weeks before impact testing as previously described. Six panels (3 having a thickness of 0.3 cm and 3 having a thickness of 0.6 cm) were tested at a drop height of 45.7 cm. All 6 panels met the test requirements. When 4 panels (2 of each thickness) were tested at the 122 cm drop height, none met the test requirements.

Examples 44 to 46

A series of security control laminates, each comprising two identical multilayer tear resistant films according to the invention, was prepared. Each multilayer film was 51 μ m thick and comprised 13 alternating layers of the stiff PET of examples 1 to 26 and 5.6 wt. % of the ductile polymeric material of the same examples. The films were coextruded onto a chilled casting wheel and were simultaneously biaxially oriented 3.3 times in each of the machine and transverse direction at 99°C. The multilayer films of examples 44 to 46 differed only in the temperature at which they were heat set. The film of example 44 was heat set at 149°C, the film of example 45 was heat set at 163°C, and the heat set temper-

ature for the film of example 46 was 177°C. A toluene/methyl ethyl ketone (T/MEK) solution (3.7:1 weight to weight ratio) of a tack free, polyester laminating adhesive was prepared by combining 6% VITEL PE-307 (commercially available from Goodyear Chemicals) and 0.3% MONDUR CB-75 (a curing agent available from Mobay Chemical Company). The solution was coated onto one of the multilayer films for each example and the coated films were dried in a circulating air oven operating at 63°C for approximately 2 minutes to provide dry adhesive coatings of approximately 0.8 g/m².

The second multilayer film for each example was laminated to the adhesive coated surface of the first multilayer film by passing the layered structures through a pair of squeeze rollers operating at 75°C and 207 dynes/cm² (30 psi). The dual film laminates were then corona treated, coated with a pressure sensitive adhesive and overcoat, and provided with a removable release liner, all as described more fully in example 43.

The panels were prepared and tested for impact resistance as described above using 0.6 cm thick glass panels and a 122 cm drop height. In example 44, 8 of the 10 panels tested met the test requirements, while 5 of the 10 panels did for example 45, and 4 of the 10 panels did for example 46. Thus, examples 44 to 46 suggest that the performance of security control laminates according to the invention can be varied by appropriate selection of the temperature at which the multilayer film is heat set. For use in security control laminates, the multilayer films of the invention are preferably heat set at about 145°C to 165°C, more preferably about 149°C.

Comparative Example 15

Comparative example 15 was a 102 µm thick commercially available security control film that comprised a pair of 51 µm thick biaxially oriented single layer PET films laminated together. Essentially, the security control laminate of comparative example 15 was like that of example 44 except that it employed single layer PET films rather than the multilayer films of the invention. Comparative example 15 was tested according to the procedure of example 43. 6 panels (3 of each thickness) evaluated at the 45.7 cm drop height met the test requirements. One panel (0.3 cm thick glass) was tested at 122 cm and met the test requirements.

Example 47

The tack free, polyester laminating adhesive of examples 44 to 46 is coated onto the multilayer film of example 44 and the coated film is dried in a circulating air oven operating 63°C for approximately 2 minutes to produce a dry adhesive coating of approximately 0.8 g/m². A second multilayer film of the same example is laminated to the adhesive coated surface of the first multilayer film by passing the layered structure through a pair of squeeze rollers operating at 75°C and 207 dynes/cm² (30 psi) to provide a dual film laminate.

A 25 µm thick biaxially oriented PET carrier film is vapor coated with aluminum to a sheet resistance of approximately 9 ohms/square using standard vapor coating techniques. The visible spectrum transmission of the aluminum coated film is approximately 18% at a wavelength of 0.55 µm. The tack free polyester laminating adhesive of example 44 is then coated onto the aluminum surface of the PET carrier film and dried to provide a dry adhesive coating of approximately 0.8 g/m². The resulting adhesive coated PET carrier film is then laminated to the uncoated surface of the dual film laminate by passing the layered structure through a pair of squeeze rollers operating at 75°C and 207 dynes/cm² (30 psi).

The exposed surface of the carrier film is corona treated as described in example 43 and a solution consisting of 100 parts hydantoin hexacrylate (HHA), 4 parts IRGACURE 184 (a photoinitiator commercially available from Ciba-Geigy Corporation), and 418 parts MEK is immediately coated onto the corona treated surface. The coated construction is then passed through a circulating air oven operating at 49°C for approximately 3 minutes to provide an HHA coating of approximately 2.7 g/m². The HHA coating is then cured by passing the construction under three banks of 200 watts/inch medium pressure mercury vapor UV lamps at a line speed of 30.5 meters/minute (100 feet/minute) and a lamp to film distance of 12 cm to provide an abrasion resistant coating.

A UV absorbent composition is prepared by combining 7.5 parts of a substituted benzophenone (e.g., UVINUL M-493 or UVINUL D-50, commercially available from BASF), 92.5 parts VITEL PE-222 (a PET terpolymer commercially available from Goodyear Chemicals), and a sufficient volume of a 1:1 weight to weight ratio T/MEK solvent system to produce a 26% solids solution.

The uncoated surface of the multilayer film laminate is corona treated to a surface energy of 40 to 44 dynes/cm (using an apparatus available from Enercon Industries) and the above described UV absorbing composition is immediately coated onto the corona treated surface. The coated laminate is then passed through a circulating air oven operating at 65°C for approximately 2 minutes to provide a UV absorbent layer having a dry coating weight of 5.4 g/m². A pressure sensitive adhesive layer is coated over the UV absorbing composition according to example 43 (where it was applied directly to the corona treated surface) to a dry coating weight of 22.6 g/m². The adhesive layer is then overcoated with a tack-free, water activatable METHOCEL layer. A 25 µm thick release liner (such as used in example 43) is removably laminated to the METHOCEL coating by passing them through a pair of squeeze rolls.

Example 48

Example 48 is similar to example 47 except that the aluminum vapor coated PET carrier film is replaced with a dyed PET film such as an optically clear dyed film commercially available from Martin Processing Company (Martinsville, VA) that is laminated to the dual film construction.

Reasonable variations and modifications are possible within the scope of the foregoing specification and drawings without departing from the invention which is defined in the accompanying claims.

Claims

1. A tear resistant film comprising at least three layers situated one on the other in a parallel array, the layers occurring essentially randomly in the array and being individually selected from a stiff polyester or copolyester and a ductile sebacic acid based copolyester; wherein the film comprises at least one layer of a stiff polyester or copolyester and at least one layer of a ductile sebacic acid based copolyester.
2. A tear resistant film according to claim 1 wherein the layers of stiff polyester or copolyester have an average nominal thickness greater than 0.5 μm .
3. A tear resistant film according to claim 1 wherein the layers of the ductile copolyester have an average nominal thickness of less than 5 μm .
4. A tear resistant film according to claim 1 wherein the layers of stiff polyester or copolyester are oriented in at least one direction.
5. A tear resistant film according to claim 1 wherein the film has a Graves area in one direction of the film equal to at least about $40 + 0.4(x)$ kpsi%, wherein x is the nominal thickness of the film in microns.
6. A tear resistant film according to claim 5 wherein the film has a tensile modulus of at least 175 kpsi in one direction of the film.
7. A tear resistant film according to claim 5 wherein the film has a Graves elongation at break of at least 20% in the tear direction of the film measured during the determination of Graves area.
8. A tear resistant film according to claim 7 wherein the film has a Graves elongation at break of at least 40% in the tear direction of the film measured during the determination of Graves area.
9. A tear resistant film according to claim 1 wherein the film comprises from about 5 to 35 layers.
10. A tear resistant film according to claim 1 wherein the stiff polyester or copolyester comprises the reaction production of (a) a dicarboxylic acid component selected from the group consisting of terephthalic acid, naphthalene dicarboxylic acid and ester derivatives thereof and (b) a diol component selected from the group consisting of ethylene glycol and 1,4-butanediol.
11. A tear resistant film according to claim 1 wherein the sebacic acid based copolyester comprises the reaction product of (a) terephthalic acid and/or naphthalene dicarboxylic acid (or ester derivatives thereof), (b) sebacic acid (or ester derivatives thereof), and (c) ethylene glycol.
12. A tear resistant film according to claim 11 comprising 20 to 80 mole equivalents terephthalic acid (or ester derivatives thereof), 80 to 20 mole equivalents sebacic acid (or ester derivatives thereof), and 100 mole equivalents ethylene glycol.
13. A tear resistant film according to claim 12 comprising 70 to 50 mole equivalents terephthalic acid (or ester derivatives thereof), correspondingly, 30 to 50 mole equivalents sebacic acid (or ester derivatives thereof), and 100 mole equivalents ethylene glycol.
14. A tear resistant film according to claim 1 wherein the sebacic acid based copolyester provides about 1 to 20 weight percent of the film.

15. A tear resistant film according to claim 1 wherein the sebacic acid based copolyester comprises the reaction product of (a) terephthalic acid and/or naphthalene dicarboxylic acid (or ester derivatives thereof), (b) sebacic acid (or ester derivatives thereof), (c) cyclohexane dicarboxylic acid (or ester derivatives thereof), and (d) ethylene glycol.
- 5 16. A tear resistant film according to claim 1 further comprising a layer of an intermediate material disposed between otherwise adjacent layers of stiff polyester or copolyester and ductile sebacic acid based copolyester.
17. A security control laminate comprising first and second tear resistant films each according to any preceding claim, the first film having a first face and a second face opposite the first face and a first layer of adhesive on the first face
10 of the first tear resistant film, the second tear resistant film being adhesively bonded to the first tear resistant film by the first layer of adhesive.
18. A security control laminate according to claim 17 further comprising a metalized layer over either the first or the second multilayer film.
- 15 19. A security control laminate according to claim 18 further comprising an abrasion resistant coating on the metalized layer.
20. A glazing unit comprising a glazing member having a first surface and a second surface opposite the first surface and a first tear resistant film according to any one of claims 1 to 16, a first face of which is adhesively secured to at least one of the first and second surfaces of the glazing member.
- 20 21. A glazing unit according to claim 20 further comprising a second tear resistant film according to claim 1 on a second face of the first tear resistant film which is opposite the first face.
- 25 22. A glazing unit according to claim 21 further comprising a metalized layer on a face of the second tear resistant film which is opposite the first tear resistant film.
- 30 23. A glazing unit according to claim 22 further comprising an abrasion resistant coating on a surface of the metalized layer opposite the second tear resistant film.

Patentansprüche

- 35 1. Reißfeste Folie, umfassend wenigstens drei Schichten, die in einer parallelen Anordnung aufeinanderliegend angeordnet sind, wobei die Schichten im wesentlichen statistisch in der Anordnung vorliegen und aus einem steifen Polyester oder Copolyester und einem duktilen Copolyester auf Sebacinsäure-Basis individuell ausgewählt sind, worin die Folie wenigstens eine Schicht eines steifen Polyesters oder Copolyesters und wenigstens eine Schicht eines duktilen Copolyesters auf Sebacinsäure-Basis umfaßt.
- 40 2. Reißfeste Folie gemäß Anspruch 1, worin die Schichten des steifen Polyesters oder Copolyesters eine durchschnittliche nominelle Dicke von mehr als 0,5 µm haben.
3. Reißfeste Folie gemäß Anspruch 1, worin die Schichten des duktilen Copolyesters eine durchschnittliche nominelle Dicke von weniger als 5 µm haben.
- 45 4. Reißfeste Folie gemäß Anspruch 1, worin die Schichten des steifen Polyesters oder Copolyesters in wenigstens einer Richtung orientiert sind.
5. Reißfeste Folie gemäß Anspruch 1, worin die Folie einen Graves-Bereich in einer Richtung der Folie hat, der
50 Wenigstens etwa $40 + 0,4(x)$ kpsi % gleich ist, worin x die nominelle Dicke der Folie in µm ist.
6. Reißfeste Folie gemäß Anspruch 5, worin die Folie einen Zugmodul von wenigstens 175 kpsi in einer Richtung der Folie hat.
- 55 7. Reißfeste Folie gemäß Anspruch 5, worin die Folie eine Graves-Reißdehnung, gemessen während der Bestimmung des Graves-Bereichs, von wenigstens 20 % in der Reißrichtung der Folie hat.
8. Reißfeste Folie gemäß Anspruch 7, worin die Folie eine Graves-Reißdehnung, gemessen während der Bestim-

mung des Graves-Bereichs, von wenigstens 40 % in der Reißrichtung der Folie hat.

9. Reißfeste Folie gemäß Anspruch 1, worin die Folie etwa 5 bis 35 Schichten umfaßt.

5 10. Reißfeste Folie gemäß Anspruch 1, worin der steife Polyester oder Copolyester das Reaktionsprodukt (a) einer Dicarbonsäure-Komponente, die aus der aus Terephthalsäure, Naphthalindicarbonsäure und Esterderivaten derselben bestehenden Gruppe ausgewählt ist, und (b) einer Diol-Komponente, die aus der aus Ethylenglycol und 1,4-Butandiol bestehenden Gruppe ausgewählt ist, umfaßt.

10 11. Reißfeste Folie gemäß Anspruch 1, worin der auf Sebacinsäure basierende Copolyester das Reaktionsprodukt (a) von Terephthalsäure und/oder Naphthalindicarbonsäure (oder Esterderivaten derselben), (b) Sebacinsäure (oder Esterderivate derselben) und (c) Ethylenglycol umfaßt.

15 12. Reißfeste Folie gemäß Anspruch 11, umfassend 20 bis 80 Mol-Äquivalente Terephthalsäure (oder Esterderivate derselben), 80 bis 20 Mol-Äquivalente Sebacinsäure (oder Esterderivate derselben) und 100 Mol-Äquivalente Ethylenglycol.

20 13. Reißfeste Folie gemäß Anspruch 12, umfassend 70 bis 50 Mol-Äquivalente Terephthalsäure (oder Esterderivate derselben), entsprechend 30 bis 50 Mol-Äquivalente Sebacinsäure (oder Esterderivate derselben) und 100 Mol-Äquivalente Ethylenglycol.

14. Reißfeste Folie gemäß Anspruch 1, worin der auf Sebacinsäure basierende Copolyester etwa 1 bis 20 Gew.-% der Folie ausmacht.

25 15. Reißfeste Folie gemäß Anspruch 1, worin der auf Sebacinsäure basierende Copolyester das Reaktionsprodukt von (a) Terephthalsäure und/oder Naphthalindicarbonsäure (oder Esterderivate derselben), (b) Sebacinsäure (oder Esterderivate derselben), (c) Cyclohexandicarbonsäure (oder Esterderivate derselben) und (d) Ethylenglycol umfaßt.

30 16. Reißfeste Folie gemäß Anspruch 1, die weiterhin eine Schicht eines intermediären Materials umfaßt, das zwischen ansonsten benachbarten Schichten aus steifem Polyester oder Copolyester und duktilem Copolyester auf Sebacinsäure-Basis angeordnet ist.

35 17. Sicherheitskontroll-Laminat, umfassend eine erste und eine zweite reißfeste Folie gemäß irgendeinem der vorhergehenden Ansprüche, wobei die erste Folie eine erste Oberfläche und eine Zweite Oberfläche gegenüber der ersten Oberfläche und eine erste Klebstoffschicht auf der ersten Oberfläche der ersten reißfesten Folie aufweist, die zweite reißfeste Folie mit der ersten reißfesten Folie durch die erste Klebstoffschicht durch Kleben verbunden ist.

40 18. Sicherheitskontroll-Laminat gemäß Anspruch 17, das weiterhin eine metallisierte Schicht entweder über der ersten oder der zweiten mehrschichtigen Folie umfaßt.

45 19. Sicherheitskontroll-Laminat gemäß Anspruch 18, das weiterhin eine abriebbeständige Beschichtung auf der metallisierten Schicht umfaßt.

50 20. Verglasungseinheit, umfassend ein Verglasungsteil mit einer ersten Oberfläche und einer zweiten Oberfläche gegenüber der ersten Oberfläche und eine erste reißfeste Folie gemäß irgendeinem der Ansprüche 1 bis 16, wobei eine erste Oberfläche an wenigstens einer der ersten und zweiten Oberfläche des Verglasungsteil durch Kleben befestigt ist.

21. Verglasungseinheit gemäß Anspruch 20, weiterhin umfassend eine zweite reißfeste Folie gemäß Anspruch 1 auf einer zweiten Oberfläche der ersten reißfesten Folie, die der ersten Oberfläche gegenüberliegt.

55 22. Verglasungseinheit gemäß Anspruch 21, weiterhin umfassend ein metallisierte Schicht auf einer Oberfläche der zweiten reißfesten Folie, die der ersten reißfesten Folie gegenüberliegt.

23. Verglasungseinheit gemäß Anspruch 22, weiterhin umfassend eine abriebbeständige Beschichtung auf einer Oberfläche der metallisierten Schicht, die der zweiten reißfesten Folie gegenüberliegt.

R revendications

- 5 1. Film résistant à la déchirure comprenant au moins trois couches placées les unes sur les autres en faisceau parallèle, les couches étant disposées de façon essentiellement aléatoire dans le faisceau et étant choisies individuellement parmi un polyester ou un copolyester rigide et un copolyester à base d'acide sébacique ductile; dans lequel le film comprend au moins une couche d'un polyester ou d'un copolyester rigide et au moins une couche d'un copolyester à base d'acide sébacique ductile.
- 10 2. Film résistant à la déchirure selon la revendication 1, dans lequel les couches de polyester ou de copolyester rigide ont une épaisseur nominale moyenne supérieure à 0,5 μm .
3. Film résistant à la déchirure selon la revendication 1, dans lequel les couches de copolyester ductile ont une épaisseur nominale moyenne inférieure à 5 μm .
- 15 4. Film résistant à la déchirure selon la revendication 1, dans lequel les couches de polyester ou de copolyester rigide sont orientées dans au moins une direction.
- 20 5. Film résistant à la déchirure selon la revendication 1, dans lequel le film a une surface de Graves dans une direction du film égale à au moins environ $40 + 0,4(x)\%$ kpsi, où x est l'épaisseur nominale du film en microns.
6. Film résistant à la déchirure selon la revendication 5, dans lequel le film possède un module de traction d'au moins 175 kpsi dans une direction du film.
- 25 7. Film résistant à la déchirure selon la revendication 5, dans lequel le film possède un allongement de Graves à la rupture d'au moins 20% dans la direction de la déchirure du film, mesuré pendant la détermination de la surface de Graves.
- 30 8. Film résistant à la déchirure selon la revendication 7, dans lequel le film possède un allongement de Graves à la rupture d'au moins 40% dans la direction de la déchirure du film, mesuré pendant la détermination de la surface de Graves.
9. Film résistant à la déchirure selon la revendication 1, dans lequel le film comprend d'environ 5 à 35 couches.
- 35 10. Film résistant à la déchirure selon la revendication 1, dans lequel le polyester ou le copolyester rigide comprend le produit de réaction (a) d'un constituant acide dicarboxylique choisi dans le groupe constitué par l'acide téréphtalique, l'acide naphtalènedicarboxylique et les dérivés esters de ceux-ci et (b) d'un constituant diol choisi dans le groupe constitué par l'éthylèneglycol et le 1,4-butanediol.
- 40 11. Film résistant à la déchirure selon la revendication 1, dans lequel le copolyester à base d'acide sébacique comprend le produit de réaction (a) de l'acide téréphtalique et/ou de l'acide naphtalènedicarboxylique (ou de dérivés esters de ceux-ci), (b) de l'acide sébacique (ou de dérivés esters de celui-ci), et (c) de l'éthylèneglycol.
- 45 12. Film résistant à la déchirure selon la revendication 11 comprenant 20 à 80 équivalents en moles d'acide téréphtalique (ou de dérivés esters de celui-ci), 80 à 20 équivalents en moles d'acide sébacique (ou de dérivés esters de celui-ci) et 100 équivalents en moles d'éthylèneglycol.
- 50 13. Film résistant à la déchirure selon la revendication 12 comprenant 70 à 50 équivalents en moles d'acide téréphtalique (ou de dérivés esters de celui-ci), de façon correspondante, 30 à 50 équivalents en moles d'acide sébacique (ou de dérivés esters de celui-ci) et 100 équivalents en moles d'éthylèneglycol.
- 55 14. Film résistant à la déchirure selon la revendication 1, dans lequel le copolyester à base d'acide sébacique fournit environ 1 à 20% en poids du film.
15. Film résistant à la déchirure selon la revendication 1, dans lequel le copolyester à base d'acide sébacique comprend le produit de réaction (a) de l'acide téréphtalique et/ou de l'acide naphtalènedicarboxylique (ou de dérivés esters de ceux-ci), (b) de l'acide sébacique (ou de dérivés esters de celui-ci), (c) de l'acide cyclohexanedicarboxylique (ou de dérivés esters de celui-ci) et (d) de l'éthylèneglycol.

16. Film résistant à la déchirure selon la revendication 1 comprenant, en outre, une couche de matière intermédiaire disposée entre des couches, par ailleurs adjacentes, de polyester ou de copolyester rigide et de copolyester à base d'acide sébacique ductile.

5 17. Stratifié de contrôle de sécurité comprenant un premier film résistant à la déchirure et un second film résistant à la déchirure, tous deux selon l'une quelconque des revendications précédentes, le premier film ayant une première face et une seconde face opposée à la première face et une première couche d'adhésif sur la première face du premier film résistant à la déchirure, le second film résistant à la déchirure étant lié par adhérence au premier film résistant à la déchirure par la première couche d'adhésif.

10 18. Stratifié de contrôle de sécurité selon la revendication 17 comprenant, en outre, une couche métallisée sur le premier film multicouche ou sur le second film multicouche.

15 19. Stratifié de contrôle de sécurité selon la revendication 18 comprenant, en outre, un revêtement résistant à l'abrasion sur la couche métallisée.

20 20. Unité de vitrage comprenant un élément de vitrage ayant une première surface et une seconde surface opposée à la première surface, et un premier film résistant à la déchirure selon l'une quelconque des revendications 1 à 16, dont une première face est fixée par adhérence au moins une de la première surface ou de la seconde surface de l'élément de vitrage.

21. Unité de vitrage selon la revendication 20 comprenant, en outre, un second film résistant à la déchirure selon la revendication 1 sur la seconde face du premier film résistant à la déchirure qui est opposée à la première face.

25 22. Unité de vitrage selon la revendication 21 comprenant, en outre, une couche métallisée sur une face du second film résistant à la déchirure qui est opposée au premier film résistant à la déchirure.

30 23. Unité de vitrage selon la revendication 22 comprenant, en outre, un revêtement résistant à l'abrasion à la surface de la couche métallisée opposée au second film résistant à la déchirure.

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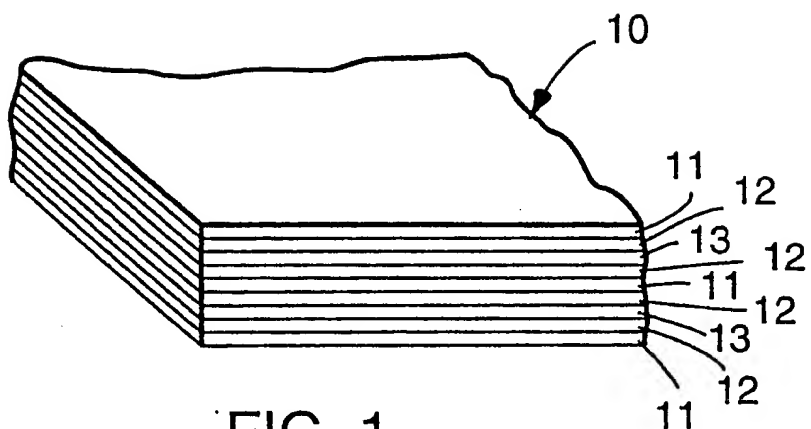


FIG. 1

FIG. 2

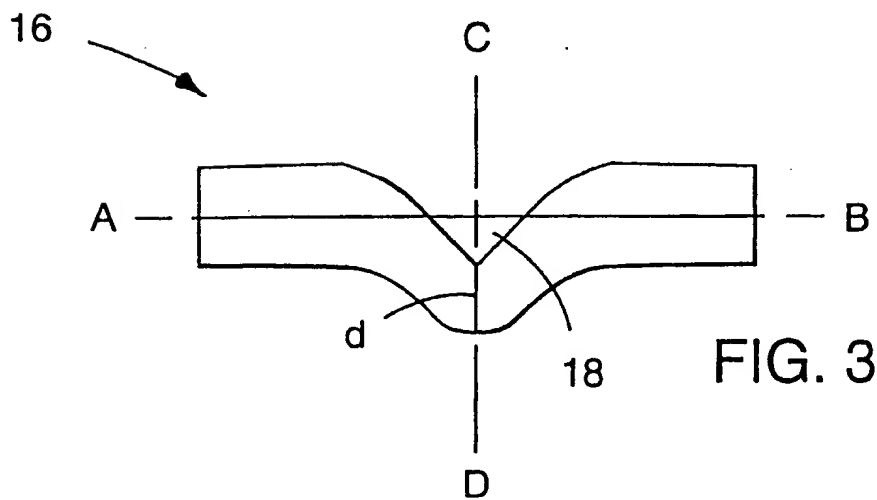
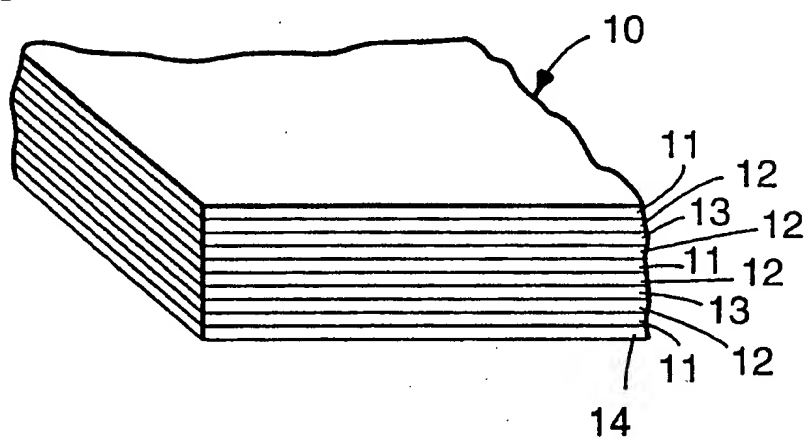
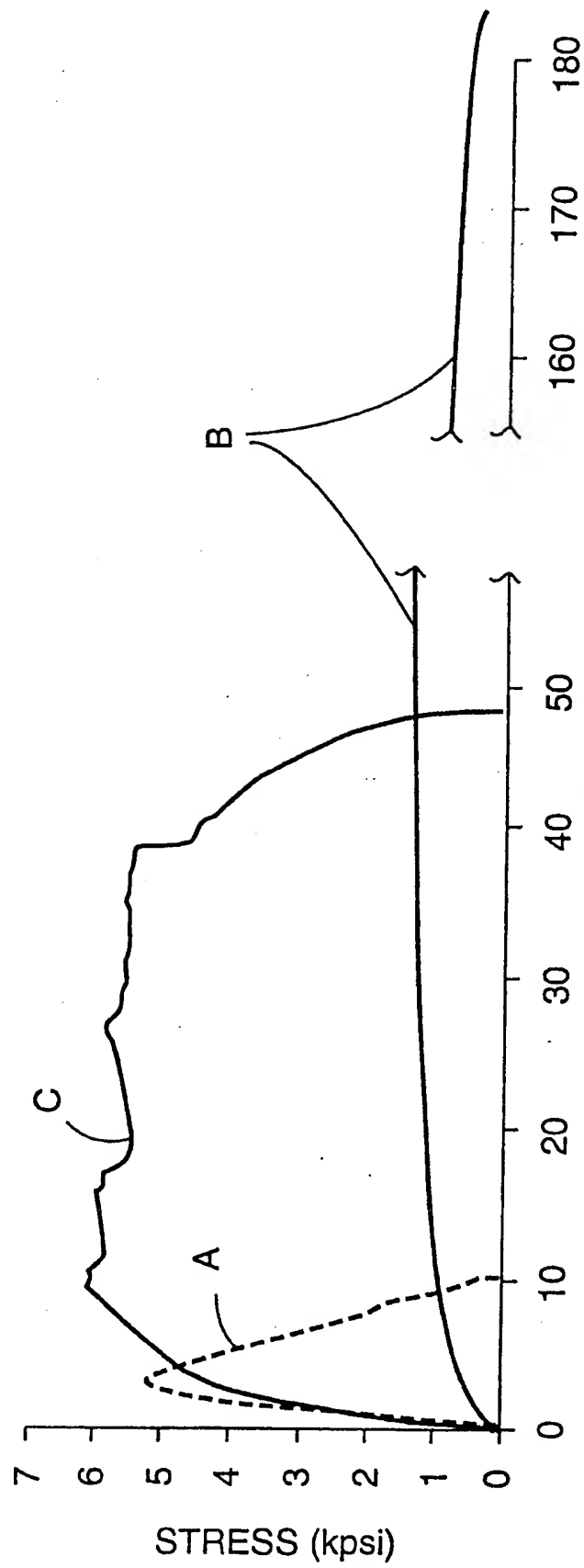


FIG. 3



GRAVES ELONGATION (%)

FIG. 4

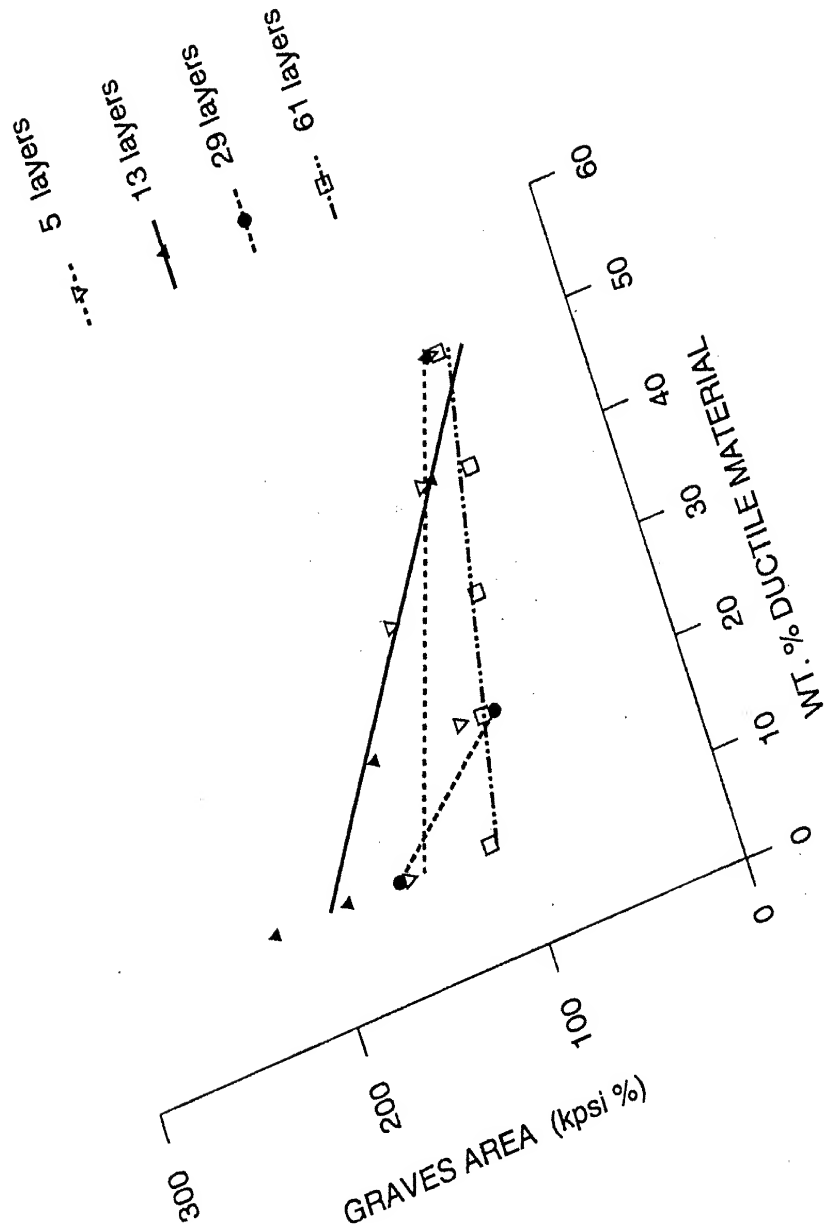


FIG. 5

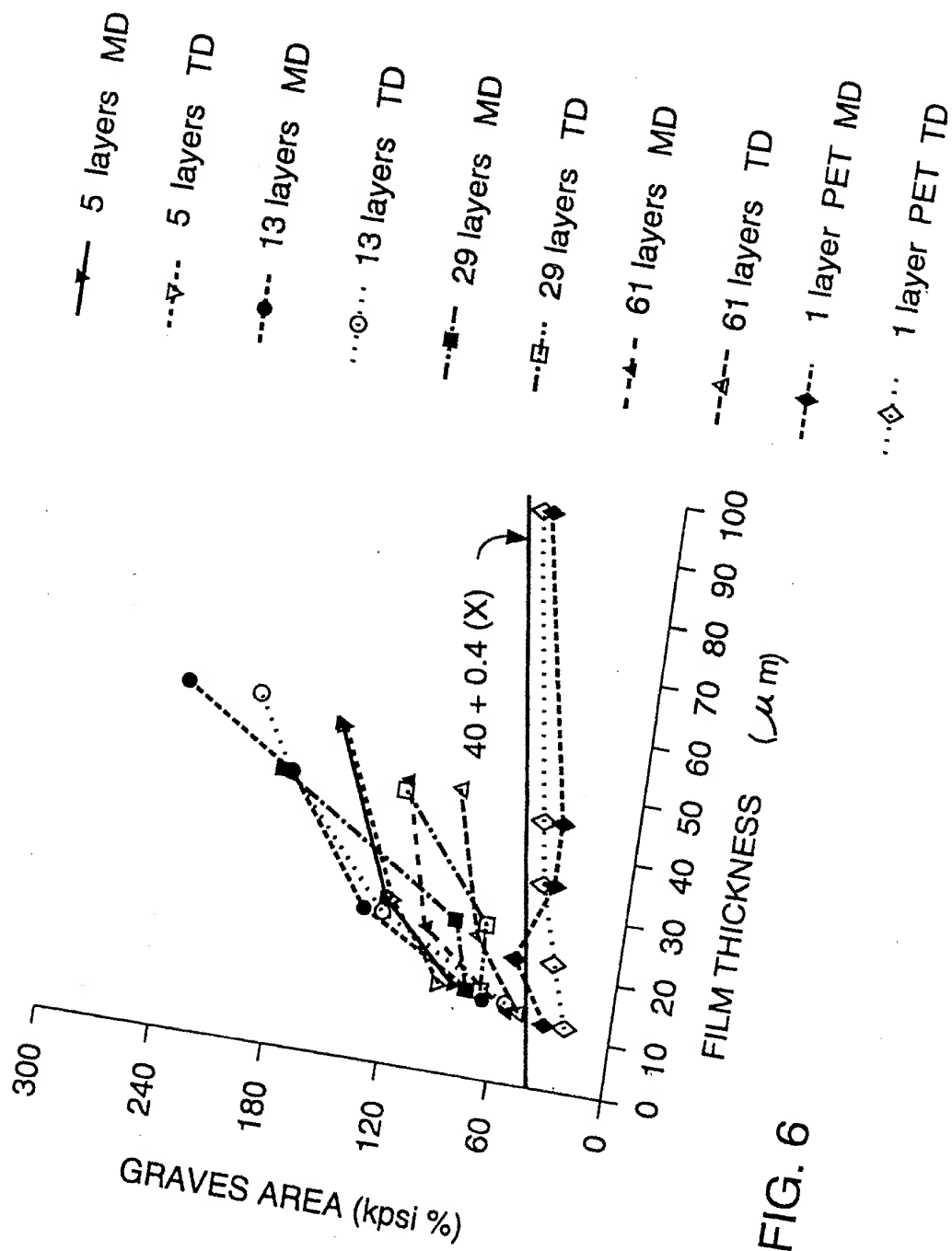


FIG. 6

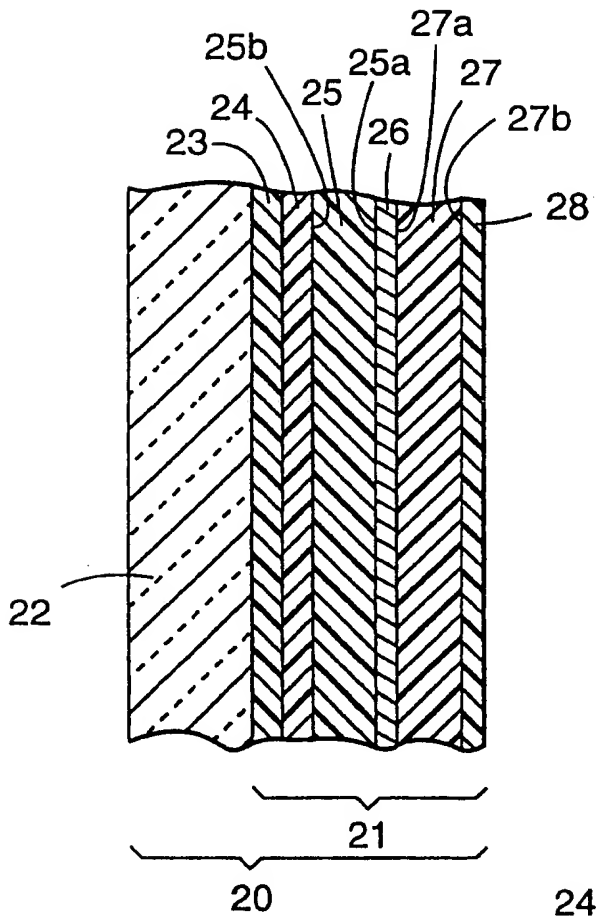
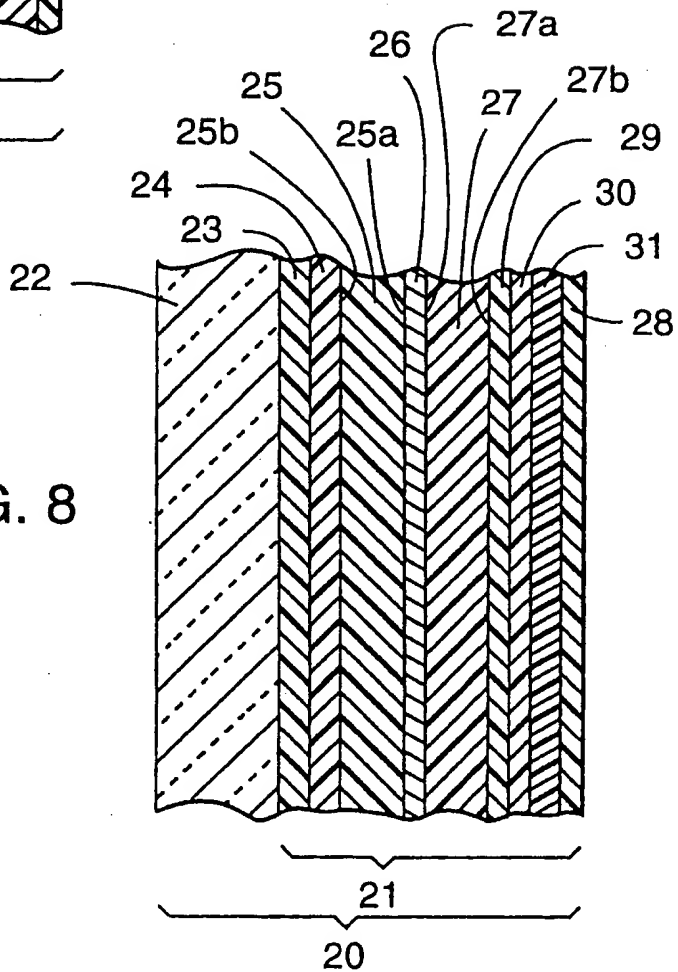


FIG. 7

FIG. 8



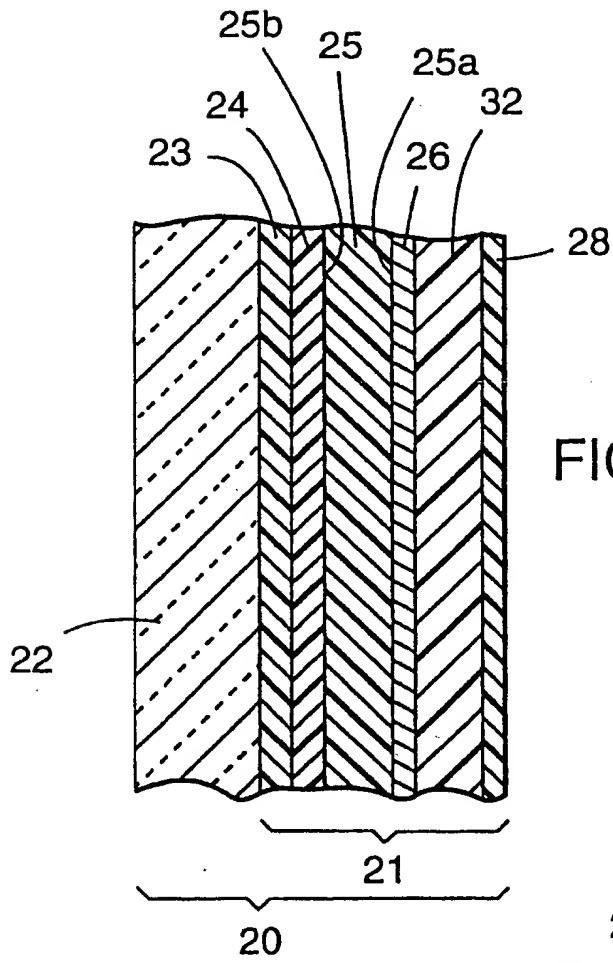


FIG. 10

